

STIC-Biotech/ChemLib

60001

From: Ramirez, Delia
Sent: Friday, February 08, 2002 7:08 PM
To: STIC-Biotech/ChemLib
Subject: case 09/606129

Hi,

I would like to request the following searches (09/606129 Maines et al.)

1. a standard search of seq id 1, 3, 18, 19, 34, 35 in the protein databases (commercial and interference)
2. an oligo search of seq id 18, 19, 34, 35 in the protein databases (commercial)

Thank you,

Delia M. Ramirez, Ph.D.
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Case 09/606129

Searcher: _____
Phone: _____
Location: *2/18/02*
Date Picked Up: *2/18/02*
Date Completed: *2/19/02*
Searcher Prep/Review: _____
Clerical: _____
Online time: _____

TYPE OF SEARCH:
NA Sequences: _____
AA Sequences: *10*
Structures: _____
Bibliographic: _____
Litigation: _____
Full text: _____
Patent Family: _____
Other: _____

VENDOR/COST (where applic.)
STN: _____
DIALOG: _____
Questel/Orbit: _____
DRLink: _____
Lexis/Nexis: _____
Sequence Sys.: *PT*
WWW/Internet: _____
Other (specify): _____

RECEIVED
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(STIC)
FBI - WASH D.C.

Pending Nucleic Acid and/or Pending Amino Acid database searches now generate two sets of results. These databases were split into two parts to reduce the time needed to update the databases daily. The split freed up more machine time for processing searches.

Searches run against the Nucleic Acid Pending database produce two sets of results, with the extensions, **.rnpm** and **.rnpn**

Searches run against the Amino Acid Pending database produce two sets of results, with the extensions, **.rapm** and **.rapn**

The Pending database search results should not be left in the case because they contain data that is confidential.

Gencore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model
Run on: February 12, 2002, 12:03:24 : Search time 94.82 Seconds
(without alignments)
10.798 Million cell updates/sec

Title: US-09-606-129A-18
Perfect score: 41
Sequence: 1 KKRIMHC 7

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 473505 seqs, 146272329 residues

Total number of hits satisfying chosen parameters: 473505

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database : SPIREMBL_17;*

```

1: sp_archea;*
2: sp_bacteria;*
3: sp_fungi;*
4: sp_human;*
5: sp_invertbrate;*
6: sp_mammal;*
7: sp_minic;*
8: sp_organelle;*
9: sp_phage;*
10: sp_plant;*
11: sp_rabbit;*
12: sp_virus;*
13: sp_vertebrate;*
14: spUnclassified;*

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Database :

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1: sp_archea;*
2: sp_bacteria;*
3: sp_fungi;*
4: sp_human;*
5: sp_invertbrate;*
6: sp_mammal;*
7: sp_minic;*
8: sp_organelle;*
9: sp_phage;*
10: sp_plant;*
11: sp_rabbit;*
12: sp_virus;*
13: sp_vertebrate;*
14: spUnclassified;*

```

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	41	100.0	599	10 Q9FHV3	Q9fhv3 arabidopsis
2	41	100.0	633	10 O49511	O49511 arabidopsis
3	38	92.7	295	11 Q9CY64	Q9cy64 mus musculi
4	38	92.7	296	4 Q9BRW8	Q9brw8 homo sapien
5	38	92.7	303	11 O9DD21	Q9dd21 mus musculi
6	37	90.2	106	5 Q9XZ45	Q9xz45 leishmania
7	36	87.3	512	10 Q9LMLO	Q9lmlo arabidopsis
8	35	85.4	508	11 O9DGS2	Q9dgs2 mus musculi
9	34	82.9	279	11 Q9QXV3	Q9qxv3 mus musculi
10	34	82.9	401	10 Q9M149	Q9m149 arabidopsis
11	34	82.9	431	10 O04613	O04613 arabidopsis
12	33	80.5	169	2 Q03949	Q03949 arabidopsis
13	33	80.5	267	2 Q9AKS0	Q9aks0 pseudomonas
14	33	80.5	495	10 Q9ST63	Q9st63 solanum tuberosum
15	33	80.5	508	10 Q80874	Q80874 arabidopsis
16	32	78.0	102	10 Q95738	Q9s738 lycopersico
17	32	78.0	104	12 Q9Q553	Q9q553 human immunodeficiency virus
18	32	78.0	108	2 P75909	P75909 escherichia coli
19	32	78.0	111	4 Q9H007	Q9h007 homo sapien

ALIGNMENTS

RESULT 1	Q9FHV3	PRELIMINARY;	PRT;	599 AA.
ID Q9FHV3;	AC Q9FHV3;			
OPRHV3;				
DT 01-MAR-2001 (TREMBLrel. 16, Created)	DT 01-MAR-2001 (TREMBLrel. 16, Last sequence update)			
DT 01-MAR-2001 (TREMBLrel. 16, Last annotation update)	DE MUTATOR-LIKE TRANSPPOSE-LIKE PROTEIN			
OS Arabidopsis thaliana (Mouse ear cress).	OS Arabidopsis thaliana (Mouse ear cress).			
OC Spermatophyta; Magnoliophyta; Embryophyta; Tracheophyta; euroids III; Brassicales; Brassicaceae; Arabidopsis.	OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae.			
NCI_TAXID=3702;				
[1]				
RP SEQUENCE FROM N.A.				
RC STRAIN-COLUMBIA;				
RX MEDLINE=94939451; Published=10470830;				
RA Kaneko T., Katoh T., Sato S., Nakamura Y., Asamizu E., Kotani H., Miyajima N., Tabata S.;				
RA "Structural analysis of Arabidopsis thaliana chromosome 5. IX. Sequence features of the regions of 1,001,550 bp covered by seventeen P1 and TAC clones."				
RT RT DNA Res. 6:183-195(1999).	RT DR EMAIL: ABO1706@BABA136.1; "			
RT DR EMAIL: ABO1706@BABA136.1; "	DR EMAIL: ABO1706@BABA136.1; "			
SEQUENCE 599 AA; 69407 MW;	SEQUENCE 599 AA; 69407 MW;			
SEQUENCE 599 AA; 69407 MW;	36BAEE2FF2AB2D717 CRC64;			
Query Match 100.0%; Score 41; DB 10; Length 599;				
Best Local Similarity 100.0%; Mismatches 0; Pred. No. 0.73;				
Matches 7; Conservative 0; Indels 0; Gaps 0;				
QY 1 KKRIMHC 7				
Db 563 KKRIMHC 569				
RESULT 2				
O49511 PRELIMINARY;				
ID O49511				
AC O49511;				
DT 01-JUN-1998 (TREMBLrel. 06, Created)	DT 01-JUN-1998 (TREMBLrel. 06, Last sequence update)			

DT	01-MAR-2001	(TREMBLrel. 16; Last annotation update)
DE	MUDR TRANSPOSSABLE ELEMENT - LIKE PROTEIN (MUDR TRANSPOSABLE ELEMENT - LIKE PROTEIN).	
DE	DE	
GN	F28J12.70 OR At4G18410.	
OS	Arabidopsis thaliana (Mouse-ear cress);	
OC	Eukaryota; Viridiplantae; Streptophytina; Embryophyta; Tracheophyta;	
OC	spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;	
OC	eurosid III; Brassicales; Brassicaceae; Arabicopsids.	
OX	NCBI_TAXID=3702;	
RN	[1]	
RP	SEQUENCE FROM N.A.	
RA	Reuven M., Willert H., Braun M., Holzer E., Brandt A., Duesterhoeft A.,	
RA	Bancroft I., Mewes H.W., Mayer K., Schueler C.,	
RA	Submitted (FEB-1998) to the EMBL/GenBank/DBJ databases.	
RL	[2]	
RN	SEQUENCE FROM N.A.	
RA	Hilbert H., Braun M., Holzer E., Brandt A., Duesterhoeft A.,	
RA	Mewes H.W., Lemke K., Mayer K.F.X.,	
RA	Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.	
RL	[3]	
RN	SEQUENCE FROM N.A.	
RA	EU Arabidopsis sequencing project;	
RA	Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.	
RL	EMBL: AL027110; CAA167211; -	
DR	EMBL: AL161548; CAB79843.1; -	
DR	Mendel; 27564; Arath13233.27564.	
DR	SEQUENCE 633 AA; 72930 MW; 3FC298BF2218C623 CRC64;	
QY	1 KKRIMHC 7	100.0%; Score 41; DB 10; Length 633;
	Best Local Similarity 100.0%; Pred. No. 0.76;	
Db	591 KKRIMHC 597	Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
RESULT	3	Query Match
Q9CY64	PRELIMINARY; PRT; 295 AA.	
AC	Q9CY64.	
ID	Q9CY64.	
DT	01-JUN-2001 (TREMBLrel. 17, Created)	
DT	01-JUN-2001 (TREMBLrel. 17, Last sequence update)	
DT	01-JUN-2001 (TREMBLrel. 17, Last annotation update)	
DE	2500001N03R1K PROTEIN.	
GN	2500001N03R1K.	
OS	Mus musculus (Mouse).	
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	
OC	Mammalia; Eutheria; Rodentia; Muridae; Murinae; Mus.	
OX	NCBI_TAXID=10090;	
RN	[1]	
RP	SEQUENCE FROM N.A.	
RC	STRAIN=C57BL/6J; TISSUE=EMBRYONIC LIVER;	
RX	MEDLINE=2157566; PubMed=1121781;	
RA	Kawai J., Shinagawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,	
RA	Schramm L.M., Staubli F., Suzuki R., Tomita M., Wagner L., Washio T.,	
RA	Sakai K., Okido T., Hara A., Furukoshi Y., Konno H., Adachi J., Fukuda S.,	
RA	Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamamoto T.,	
RA	Saito T., Okazaki Y., Gojobori T., Bono H., Kasukawa T., Saito R.,	
RA	Kadota K., Matsuo H.A., Ashburner M., Batzalov S., Casavant T.,	
RA	Fleischmann W., Gaasterland T., Giess C., King B., Kochiwa H.,	
RA	Kueni P., Lewis S., Matsuo Y., Nikaido T., Pesole G., Quackenbush J.,	
RA	Schramm L.M., Marchionni L., Mashima J., Mazzarelli J., Rodriguez J.,	
RA	Nordone P., Ringwald M., Schoenbach C., Sakamoto N., Storch K.-P.,	
RA	Sasaki H., Sato K., Seiya T., Shibusawa Y., Whitaker C., Wilming L.,	
RA	Suzuki H., Toyo-oka K., Wang K.H., Weitz C., Hayashizawa-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kontsuki S.,	
RA	Hayashizawa-Kurihara K., Hasegawa Y., Kawaji H., Kontsuki S.,	

"Functional annotation of a full-length mouse cDNA collection." RT
 Nature 409:655-690(2001).
 RL EMBL; Ak01084; BB27219; 1;
 DR MG/D; MGI:1917355; 250001IN0RHK.
 DR InterPro: IPI000683; GRO_IDH_MoCA.
 DR Pfam: PF01408; GFO_IDH_MoCA; 1.
 DR SEQUENCE: F01408; F2E1682BD77032A4 CRC64;
 SQ 295 AA; 33524 MW;

Query Match 92.7%; Score 38; DB 11; Length 295;
 Best Local Similarity 85.7%; Pred. No. 1.8;
 Matches 6; Conservative 1; Mismatches 0; Indels 0;
 Qy: 1 KKRIMHC 7
 Db 274 KKRILHC 280

RESULT 4
 Q9BRW8 PRELIMINARY; PRT; 296 AA.
 ID Q9BRW8; PRELIMINARY; PRT; 296 AA.
 AC Q9BRW8;
 DT 01-JUN-2001 (TREMBLrel. 17, Created)
 DT 01-JUN-2001 (TREMBLrel. 17, Last sequence update)
 DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
 DE SIMILAR TO BIOLIVER IN REDUCTASE A.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo
 OC NCBI_TAXID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=BRAIN;
 RA Strausberg R.;
 RL Submitted (APR-2001) to the EMBL/GenBank/DDBJ databases.
 DR EMBL; BC0590; AAH0590; 1;
 SQ SEQUENCE 296 AA; 33432 MW; A389AFDBB6ADBAFF CRC64;

Query Match 92.7%; Score 38; DB 4; Length 296;
 Best Local Similarity 85.7%; Pred. No. 1.8;
 Matches 6; Conservative 1; Mismatches 0; Indels 0;
 Qy: 1 KKRIMHC 7
 Db 275 KKRILHC 281

RESULT 5
 Q9DD21 PRELIMINARY; PRT; 303 AA.
 ID Q9DD21; PRELIMINARY; PRT; 303 AA.
 AC Q9DD21;
 DT 01-JUN-2001 (TREMBLrel. 17, Created)
 DT 01-JUN-2001 (TREMBLrel. 17, Last sequence update)
 DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
 DE 0610006A1RIK PROTEIN.
 GN 0610006A1RIK
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mu
 NCBI_TAXID=10990;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=KIDNEY;
 RX MEDLINE=21085560; PubMed=11217851;
 RA Arakawa T., Shiranaga A., Shibata K., Yoshino M., Itoh M., Ishii Y,
 RA Arakawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaoka I,
 RA Saito T., Okada Y., Gotoh T., Bono H., Kasukawa T., Saito R,
 RA Kadoya K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,
 RA Fleischmann W., Gaasterland T., Gissi C., King B., Kochiwa H.,
 RA Kuehl P., Lewis S., Matsuo Y., Niikido I., Pesole G., Quackenbush F.,
 RA Schramm L.M., Staubli F., Suzuki R., Tomita M., Wagner L., Washi

RA	Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G., Blieke J., Boffiglia D., Bojunga N., Carninci P., de Bonaldo M.F., Brownstein M.J., Built C., Fletcher C., Fujita M., Gariboldi M., Gustincich S., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H., Lyons P., Marchionni L., Mashima J., Mazzarelli P., Nodone P., Nordin J., Rodriguez I., Samamoto N., Saeki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F., Suzuki H., Toyo-oka K., Wang K.H., Weitz C., Whittaker C., Wilming L., Wyshaw-Boris A., Yoshida K., Kawaji H., Kohtsuka S., Hayashizaki Y.; "Functional annotation of a full-length mouse cDNA collection.";	RA	P10K1_10.
RA	Nature 409:655-690(2001).	RA	Arabidopsis thaliana (Mouse-ear cress).
RA	EMBL; AK002231; BAB21900_1; MGII; MGII:191150; 0610006A1Rik.	RA	OS
RA	InterPro; IPRO000684; GFO_IDH_Moca.	RA	OC
RA	Pfam; PF01408; GFO_IDH_Moca; 1.	RA	SPERMATOPHYTA; Magnoliophyta; eudicotyledons; core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae; Arabicopsis.
RA	SEQUENCE 303 AA; 34491 MW; 52D8A3B02EE956EB CRC64;	RA	NCBI_TAXID=702;
RT	[1]	RN	OX
RT	SEQUENCE FROM N.A.	RP	SEQUENCE FROM N.A.
RT	STRAIN=CV_COLUMBIA;	RC	STRAIN=CV_COLUMBIA;
RT	Liu S.X., Chan A., Yu G., Etgu P., Lee J.M., Lenz C., Pham P., Sekano H., Toriumi M., Chung M., Goldsmith A., Liu A., Smith A., Vaysberg M., Altafi H., Brooks S., Buehler E., Chao Q., Conn L., Conway A., Hansen N., Johnson-Hopson C., Khan S., Kim C., Lam B., Miranda M., Nguyen M., Palm C.J., Shinn P., Southwick A., Davis R.W., Ecker J.R., Federspiel N.A., Theologis A., "The sequence of BAC P10K1 from Arabidopsis thaliana chromosome 1.;"	RA	RA
RT	[2]	RN	RA
RT	Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.	RA	RA
RT	SEQUENCE FROM N.A.	RC	SEQUENCE FROM N.A.
RT	STRAIN=CV_COLUMBIA;	RA	SEQUENCE FROM N.A.
RT	Theologis A.;	RA	Theologis A.;
RT	Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.	RL	Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
RT	[3]	RN	RA
RT	SEQUENCE FROM N.A.	RP	SEQUENCE FROM N.A.
RT	STRAIN=CV_COLUMBIA;	RC	SEQUENCE FROM N.A.
RT	Theologis A.;	RA	Theologis A.;
RT	Submitted (JUN-2000) to the EMBL/GenBank/DBJ databases.	RL	Submitted (JUN-2000) to the EMBL/GenBank/DBJ databases.
RT	[4]	RN	RA
RT	SEQUENCE FROM N.A.	RP	SEQUENCE FROM N.A.
RT	STRAIN=CV_COLUMBIA;	RC	SEQUENCE FROM N.A.
RT	Theologis A.;	RA	Theologis A.;
RT	Submitted (JUL-2000) to the EMBL/GenBank/DBJ databases.	RL	Submitted (JUL-2000) to the EMBL/GenBank/DBJ databases.
RT	[5]	RN	RA
RT	CORACTOR: FAD (BY SIMILARITY);	CC	CORACTOR: FAD (BY SIMILARITY);
RT	-!- SIMILARITY: TO PYRIDINE NUCLEOTIDE-DISULPHIDE OXIDOREDUCTASES CLASS-II.	CC	-!- SIMILARITY: TO PYRIDINE NUCLEOTIDE-DISULPHIDE OXIDOREDUCTASES CLASS-II.
RT	EMBL; AC067971; AAFB2202_1; -	DR	EMBL; AC067971; AAFB2202_1; -
RT	InterPro; IPR01327; P450_reddox.	DR	InterPro; IPR01327; P450_reddox.
RT	Pfam; PF00070; P450_reddox.	DR	Pfam; PF00070; P450_reddox.
RT	FAD: Flavoprotein; Oxidoreductase; Redox-active center.	KW	FAD: Flavoprotein; Oxidoreductase; Redox-active center.
RT	SEQUENCE 512 AA; 56857 MW; 1F63AF9A1A2C13B CRC4;	SQ	SEQUENCE 512 AA; 56857 MW; 1F63AF9A1A2C13B CRC4;
RT	[6]	RN	RA
RT	Query Match 92.7%; Score 38; DB 11; Length 303; Best Local Similarity 85.7%; Pred. No. 1.8; Mismatches 0; Indels 0; Gaps 0;	Qy	Query Match 97.8%; Score 36; DB 10; Length 512; Best Local Similarity 71.4%; Pred. No. 7.5; Mismatches 0; Indels 0; Gaps 0;
RT	Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;	Db	Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
RT	[7]	Q9XNM5	Q9XNM5
RT	PRELIMINARY; PRY; 106 AA.	PR	PRELIMINARY; PRY; 106 AA.
RT	[8]	Q9XNM5	Q9XNM5
RT	AC	AC	AC
RT	01-NOV-1999 (TREMBLrel. 12, Created)	DT	01-JUN-2001 (TREMBLrel. 17, Last sequence update)
RT	01-MAY-2000 (TREMBLrel. 13, Last sequence update)	DT	01-JUN-2001 (TREMBLrel. 17, Last annotation update)
RT	01-JUN-2001 (TREMBLrel. 17, Last annotation update)	DT	01-JUN-2001 (TREMBLrel. 17, Last annotation update)
RT	RIBOSOMAL PROTEIN L44.	DE	RIBOSOMAL PROTEIN L44.
RT	RPL4.	DR	RPL4.
RT	Leishmania amazonensis.	OS	Leishmania amazonensis.
RT	Eukaryotes; Euglenozoa; Kinetoplastida; Trypanosomatidae; Leishmanida.	OC	Eukaryotes; Euglenozoa; Kinetoplastida; Trypanosomatidae; Leishmanida.
RT	NCBL_TAXID=5659;	NCBL_TAXID=5659;	NCBL_TAXID=5659;
RT	[9]	RN	RN
RT	SEQUENCE FROM N.A.	RP	SEQUENCE FROM N.A.
RT	STRAIN=Lv18;	RC	STRAIN=Lv18;
RT	Porter-Kelley J., Chaudhuri G.; "Cloning and characterization of Leishmania ribosomal protein L44.";	RA	Porter-Kelley J., Chaudhuri G.; "Cloning and characterization of Leishmania ribosomal protein L44.";
RT	Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.	RL	Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
RT	-!- SIMILARITY: TO THE L44E FAMILY OF RIBOSOMAL PROTEINS.	CC	-!- SIMILARITY: TO THE L44E FAMILY OF RIBOSOMAL PROTEINS.
RT	EMBL; AF148853; AA031928_2; -	DR	EMBL; AF148853; AA031928_2; -
RT	InterPro; IPRO00552; Ribosomal_L44_E.	DR	InterPro; IPRO00552; Ribosomal_L44_E.
RT	Pfam; PF00935; Ribosomal_L44_E.	DR	Pfam; PF00935; Ribosomal_L44_E.
RT	PROSITE; PS002841; Ribosomal_L44_E; 1.	DR	PROSITE; PS002841; Ribosomal_L44_E; 1.
RT	PROSITE; PS01112; RIBOSOMAL_L44_E; 1.	DR	PROSITE; PS01112; RIBOSOMAL_L44_E; 1.
RT	Ribosomal protein.	KW	Ribosomal protein.
RT	SEQUENCE 106 AA; 12283 MW; F30A3AB2047B0334 CRC64;	SQ	SEQUENCE 106 AA; 12283 MW; F30A3AB2047B0334 CRC64;
RT	[10]	RN	RN
RT	RESULT 6	Q9LML0	RESULT 7
RT	Q9LML0 PRELIMINARY; PRY; 512 AA.	ID	Q9LML0 PRELIMINARY; PRY; 512 AA.
RT	Q9LML0 ID	Q9LML0	Q9LML0 ID
RT	AC	AC	AC
RT	01-OCT-2000 (TREMBLrel. 15, Created)	DT	01-OCT-2000 (TREMBLrel. 15, Last sequence update)
RT	01-OCT-2000 (TREMBLrel. 15, Last sequence update)	DT	01-OCT-2000 (TREMBLrel. 15, Last sequence update)
RT	01-JUN-2001 (TREMBLrel. 17, Last annotation update)	DT	01-JUN-2001 (TREMBLrel. 17, Last annotation update)
RT	01-JUN-2001 (TREMBLrel. 17, Last annotation update)	DT	01-JUN-2001 (TREMBLrel. 17, Last annotation update)
RT	F10K1_10 PROTEIN	DR	F10K1_10 PROTEIN
RT	[11]	RN	RN
RT	SEQUENCE FROM N.A.	RP	SEQUENCE FROM N.A.
RT	STRAIN=CV7BL6J; TISSUE=TONGUE;	RC	STRAIN=CV7BL6J; TISSUE=TONGUE;
RT	MEDLINE=2108560; PubMed=11217851;	RX	MEDLINE=2108560; PubMed=11217851;
RT	Kawai J., Shinagawa A., Shibusawa K., Yoshino M., Itoh M., Ishii Y., Arakawa T., Hara A., Fukunaga Y., Konno H., Adachi J., Fukuda S., Aizawa K., Izawa M., Nishi K., Kiyoysawa H., Kondo S., Yamakawa T., Saito T., Okano Y., Gojobori T., Bono H., Kasukawa T., Saito T., Kadoya K., Matsuda H.A., Batalov S., Casavant T.,	RA	RA
RT	Eukaryotes; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Murinae; Murinae; Mus; NCBI_TAXID=10090;	OC	OC

RA Fleischmann W., Gaasterland T., Gissi C., King B., Kochiwa H., Lewis S., Matsuo Y., Nikaido I., Pasole G., Quackenbush J., Schriml L.M., Staubli F., Suzuki R., Tonita M., Wagner L., Washio T., Xie H., Zhou J.	Produced by alternative splicing. Tissue specificity: in the adult, widely expressed with highest levels at thymus and testis, expressed throughout the whole embryo at all stages of development examined. At day 10, highest expression is found in the yolk sac while at day 16 and 18, higher levels are found in inner compartments of bone.
RA Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G., Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F., Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M., Gustincich S., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H., Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P., Nordone P., Ring B., Ringwald M., Rodriguez J., Sakamoto N., Suzuki H., Sato K., Schoenbach C., Seya T., Shibusawa Y., Storch K.-F., Suzuki H., Toyooka K., Wang K.H., Weitz C., Whittaker C., Wilming L., Wyshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohtsuki S., Hayashizaki Y.;	-! Developmental stage: in the embryo, highest expression of isoform 1 is found at day 11 while highest expression of isoform 2 is found at day 7.
RT Functional annotation of a full-length mouse cDNA collection. /	-! Similarity: contains 1 PHD-finger domain.
RL Nature 409:685-690(2001).	DR EMBL; AF17753; AA16911; 1; DR EMBL; AF17755; AA16908; 1; DR EMBL; AF17756; AA16909; 1; DR EMBL; AF17757; AA16910; 1; DR EMBL; AF149820; AA09183; 1;
DR MBD; AR01034; BAB26655; 1; -.	DR MGII; 349481; Ingl1; DR InterPro; IPR001965; PhD.
DR MGI:1344390; Oasi.	DR Pfam; PF00628; PhD; 1;
DR InterPro; IPR01797; 25A-synth.	DR SMART; SM00249; PhD; 1;
DR InterPro; IPR01201; PAP_5A_core.	DR Oncogene; Anti-oncogene; Alternative splicing.
DR InterPro; IPR00626; Ubiquitin.	DR DOMAIN; 210 259 FT VARSPLIC 1 94 MISSING (IN ISOFORM 2). FT CONFFLICT 203 203 L->F (IN REF. 2). SQ SEQUENCE 279 AA; 32109 MW; 6765C984EEF179F4 CRC64;
DR SMART; SM00013; UBQ; 1.	Query Match 82.9%; Score 34; DB 11; Length 279; Best Local Similarity 57.1%; Pred. No. 12; Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
DR PROSITE; PS00833; 25A-SYNTH_2; UNKNOWN_1.	QY 1 KKRRIMHC 7 Db 67 KRRVILHC 73
DR PROSITE; PS01052; 25A-SYNTH_3; 1.	RESULT 10 Q9M149 PRELIMINARY; PRT; 401 AA. ID Q9M149 AC Q9M149; DT 01-OCT-2000 (TREMBLrel. 15, Created) DT 01-JUN-2001 (TREMBLrel. 17, Last sequence update) DE PUTATIVE PHOSPHATIDYLINOSITOL KINASE. GN AT4G01190 OS Arabidopsis thaliana (Mouse-ear cress). OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophytai; Magnoliophyta; euicotsyledons; core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis. OX NCBI_TaxID=3702;
DR PROSITE; PS00053; UBQUITIN_N_2; 1.	RP SEQUENCE FROM N.A. RA Lamar B.; Stoneking T.; Stumpf J.; Mewes H.W.; Lemcke K.; Mayer K.F.X.; RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases. [2] RN SEQUENCE FROM N.A. RA EU Arabidopsis sequencing project; RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases. DR AL161491; CAB80928; 1; DR InterPro; IPR002498; PIP5K. DR Pfam; PF01504; PIPPK; 1; DR SMART; SM00330; PIFKC; 1. KW Kinase. SQ SEQUENCE 401 AA; 45658 MW; 8A12D10DA2DED4CA CRC64;
SEQUENCE FROM N.A. AND CHARACTERIZATION.	Query Match 82.9%; Score 34; DB 10; Length 401; Best Local Similarity 85.7%; Pred. No. 16; Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
RT STRAIN=129/SVJ; TISSUE=BRAIN, EMBRYONIC FIBROBLAST, AND SPLEEN; RX MEDLINE=20011419; PubMed=10542254;	QY 1 KKRRIMHC 7 Db 353 KKRRIMHC 359
RA Zeremski M.; Hill J.E.; Kwek S.S.S.; Grigorjan I.A.; Gurova K.V.; Garkavtsev I.V.; Diatchenko L.; Koonin E.V.; Gudkov A.V.; RT Structure and regulation of the mouse ingr gene. Three alternative transcripts encode two PHD finger proteins that have opposite effects on P53 function. /	CC ACTIVATION AND MAY FUNCTION AS AN ONCOPROTEIN. ISOFORM 2 ACTS AS A NEGATIVE GROWTH REGULATOR BY COOPERATING WITH P53 IN TRANSCRIPTIONAL ACTIVATION OF P53-RESPONSIVE GENES AND MAY ACT AS A TUMOR SUPPRESSOR. CC SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY). CC ALTERNATIVE PRODUCTS: 2 ISOFORMS; 1 (SHOWN HERB) AND 2; ARE

Qy 1 KKRIMHC 7
 | :|:||
 Db 224 KRRLHc 230

RESULT 15
 080874 PRELIMINARY; PRT: 508 AA.
 ID 080874;
 AC 080874;
 DT 01-NOV-1998 (TREMBrel, 08, Created)
 DT 01-NOV-1998 (TREMBrel, 08, Last sequence update)
 DT 01-JUN-2001 (TREMBrel, 17, Last annotation update)
 DE PUTATIVE UBIQUINONE REDUCTASE.
 GN F23E1.9.
 OS Arabidopsis thaliana (Mouse-ear cress).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophytina; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
 OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
 OX NCBI_TAXID=3702;
 RN [1]
 RP SEQUENCE FROM N.A.
 STRAIN=CV COLUMBIA;
 RC Rounsley S.D., Lin X., Ketchum K.A., Crosby M.L., Brandon R.C.,
 RA Sykes S.M., Kaul S., Mason T.M., Kerlavage A.R., Adams M.D.,
 RA Somerville C.R., Venter J.C.;
 RT "Arabidopsis thaliana chromosome II BAC F23E1 genomic sequence.";
 RL Submitted (Aug-1998) to the EMBL/GenBank/DBJ databases.
 CC -|- COFACTOR: FAD (BY SIMILARITY).
 CC -|- SIMILARITY: TO PYRIDINE NUCLEOTIDE-DISULPHIDE OXIDOREDUCTASES
 CLASS-I.
 EMBL; AAC04680; AAC31853.1;
 DR InterPro: IPR001327; FAD_pyr_redox.
 DR Pfam: PF00070; PYR_FEDDX; 1.
 DR Redox; Oxidoreductase; Redox-active center.
 KW Flavoprotein; MW: 266A434E702A0C27 CRC64;
 SQ SEQUENCE 508 AA; 5603 MW;

Query Match 80.5%; Score 33; DB 10; Length 508;
 Best Local Similarity 57.1%; Pred. No. 33;
 Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 KKRIMHC 7
 | :|:||
 Db 236 KRRLHc 242

Search completed: February 12, 2002, 12:03:25
 Job time: 816 sec

Copyright (c) 1993 - 2000 Compugen Ltd.	GenCore version 4.5						
protein - protein search, using sw model							
on: February 12, 2002, 12:04:01 ; Search time 30.28 Seconds (without alignments)	gapop 10.0 , Gapext 0.5						
file: US-09-506-129A-18 score: 41 length: 1 KKRIMHC 7	BLOSUM62 gapop 10.0 , Gapext 0.5						
searched: 100059 seqs, 36664827 residues	al number of hits satisfying chosen parameters: 100059						
Maximum DB seq length: 0							
Maximum DB seq length: 200000000							
--processing: Minimum Match 0%							
Maximum Match 100%							
Listing first 45 summaries							
SwissProt_39;*							
Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.							
SUMMARIES							
8	Query	Match	Length	DB	ID	Description	
1	41	100.0	295	1	BIEA_RAT	P46844 ratus norv	
2	38	92.7	296	1	BIEA_HUMAN	P53004 homo sapien	
3	33	85.4	473	1	OASL_MOUSE	P29452 mus musculus	
4	33	80.5	925	1	PIPL_YEAST	P00220 saccharomyces cerevisiae	
5	32	78.0	140	1	EKB2_MOUSE	P45818 mus musculus	
6	32	78.0	140	1	EKB2_HUMAN	P26885 homo sapiens	
7	32	78.0	328	1	HYBA_ECOLI	P37179 escherichia coli	
8	32	78.0	528	1	DYR2_HUMAN	Q92630 homo sapiens	
9	32	78.0	571	1	AMP1_LYSESIS	Q10112 lyticus	
10	32	78.0	573	1	AMPL_SOLTU	P31427 solanum tuberosum	
11	31	75.6	105	1	RL44_TRYPB	P17743 trypansoma brucei	
12	31	75.6	105	1	Y091_NPVOB	O10341 organelle	
13	31	75.6	323	1	CYCH_XENLA	P51947 xenopus laevis	
14	31	75.6	451	1	SUN_HAEM	P44788 haemophilus ducreyi	
15	31	75.6	768	1	CUL3_HUMAN	Q13618 homo sapiens	
16	31	75.6	768	1	CUL3_MOUSE	P99115 mus musculus	
17	31	75.6	861	1	UL52_HSV7J	P52468 human herpes virus	
18	30	73.2	108	1	YZ09_METJJA	Q60221 methanococcus	
19	30	73.2	4128	1	PRKD_MOUSE	P97313 mus musculus	
20	29	70.7	38	1	RL36_THMENA	Q9X116 thermotoga maritima	
21	29	70.7	142	1	SEC6_BUCA1	P57161 buchnera aphidis	
22	29	70.7	144	1	YHP5_YEAST	P38808 saccharomyces cerevisiae	
23	29	70.7	257	1	YZG1_CAEEL	P53316 caenorhabditis elegans	
24	29	70.7	300	1	RANT_BPP22	P03037 bacteriophaga	
25	29	70.7	359	1	ODPB_AT	P49432 rat	
26	29	70.7	423	1	SHP1_YEST	P34223 saccharomyces cerevisiae	
27	29	70.7	560	1	YD2H_SCHEPO	Q10264 schizosaccharomyces pombe	
28	29	70.7	588	1	DYR3_HUMAN	Q43781 homo sapiens	
29	29	70.7	632	1	AUBF_HAEIN	P44804 haemophilus influenzae	
30	29	70.7	670	1	REP_HAEIN	P34341 ceponthobacter	
31	29	70.7	918	1	YK62_CAEEL	P24120 dirosophila melanogaster	
32	29	70.7	1059	1	CAPU_DROME	P08621 americanus	
33	29	70.7	1099	1	SEPA_EMENI	P39740 bacillus subtilis	
					ALIGNMENTS		
					RESULT 1		
					BIEA_RAT	STANDARD; PRT; 295 AA.	
					ID BIEA_RAT		
					AC P46444;	Rel. 32, Created)	
					DT 01-NOV-1995	Rel. 32, Last sequence update)	
					DT 01-NOV-1995	Rel. 32, Last annotation update)	
					DT 15-JUL-1999	Rel. 38, Last annotation update)	
					DE BILIVERDIN REDUCTASE A PRECURSOR (EC 1.3.1.24) (BILIVERDIN-IX ALPHA-REDUCTASE).		
					GN BLVR.		
					OS Rattus norvegicus (Rat).		
					OC Euarystoa; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.		
					NCBI_TaxID=10116;		
					OX		
					RN [1]	SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.	
					RC TISSUE=Kidney;		
					RX MEDLINE=92156147; PubMed=1371282;		
					RA Pakhrai H.; Maines M.D.;	"Expression and characterization of a cDNA for rat kidney biliverdin reductase. Evidence suggesting the liver and kidney enzymes are the same transcript product.";	
					RT J. Biol. Chem. 267:4023-4029(1992).		
					RL RN [2]	MUTAGENESIS.	
					RP RX	McCoubrey W.K. Jr.; Maines M.D.;	
					RA RA	"Site-directed mutagenesis of cysteine residues in biliverdin reductase. Roles in substrate and cofactor binding.";	
					RT RL Eur. J. Biochem. 222:577-603(1994).		
					CC CC	"FUNCTION: CONVERTS BILIVERDIN TO BILIRUBIN: DISPLAYS TWO DISTINCT PH OPTIMA USING A DIFFERENT COFACTOR AT EACH PH: NADH AT THE LOWER PH 6.7-6.9 RANGE AND NADPH AT PH 8.5-8.7. NADPH, HOWEVER, IS THE PROBABLE COFACTOR IN BIOLOGICAL SYSTEMS."	
					CC CC	"-1 COFACTOR: BINDS ONE ZINC ION"	
					CC CC	"-1 PATHWAY: FINAL STEP IN HEME METABOLISM."	
					CC CC	"-1 SUBUNIT: MONOMER (BY SIMILARITY)."	
					CC CC	"-1 SUBCELLULAR LOCATION: CYTOPLASMIC."	
					CC CC	"-1 SIMILARITY: TO E.COLI YHHX."	
					CC CC	This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation in the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See http://www.isb-sib.ch/announce or send an email to license@isb-sib.ch).	
					DR EMBL; M01681; AAA08303; InterPro; IPRO00683; GFO_IDH_MoCA; PFam; PF14048; GFO_IDH_MoCA; 1.		
					DR KW Oxidoreductase; NAD; NADP; Zinc.		
					FT PROPEP 1		
					FT DORNAIN 2		
					FT DORNAIN 3	295	
					FT DORNAIN 11	196	
					FT BILIVERDIN REDUCTASE A. 1		
					FT BILIVERDIN REDUCTASE A. 2		
					FT BILIVERDIN REDUCTASE A. 3		
					FT BILIVERDIN REDUCTASE A. 11		

-!- SIMILARITY: CONTAINS 1 UBIQUITIN-LIKE DOMAIN. HUMAN OASL.
 CC -!- CAUTION: THIS MAY NOT BE THE TRUE ORTHOLOG OF HUMAN OASL.

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CC EMBL: AF058835; AAD02818; 1;
 DR MGII: MGI:1344380; CasI;
 DR InterPro: IPR001797; 25A-synth.
 DR InterPro: IPR001201; PAP-25A-core.
 DR InterPro: IPR000626; Ubiquitin.
 DR Pfam: PF00240; ubiquitin_1.
 DR PROSITE: PS00832; 25A_SYNTH_1; FALSE_NEG.
 DR PROSITE: PS00833; 25A_SYNTH_2; 1.
 DR PROSITE: PS50152; 25A_SYNTH_3; 1.
 DR PROSITE: PS50053; UBIQUITIN_2; FALSE_NEG.
 KW RNA-binding; Transferase; Nucleotidyltransferase.
 FT DOMAIN: 435..473 UBIQUITIN-LIKE
 SQ SEQUENCE 473 AA; 54625 MW; 570E0E08A51C8460 CRC64;

RESULT 4
 ID PIP1_YEAST STANDARD; PRT; 925 AA.
 AC P40020;
 DT 01-FEB-1995 (Rel. 31; Created)
 DT 01-NOV-1997 (Rel. 35; Last sequence update)
 DE POLYMERASE-INTERACTING PROTEIN 1 (FACTOR INTERACTING WITH REF)
 GN PIP1 OR FIR1 OR YER032W.
 OS Saccharomyces cerevisiae (Baker's yeast).
 OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomyces.
 OX NCBI_TAXID=4932;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=W303;
 RA del Olmo M., Gross S., Moore C.L.; Submitted (FEB-1997) to the EMBL/GenBank/DDBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=S288C / AB972;
 RA Aviles E., Berno A., Brennan T., Carpenter J., Chen E., Cherry J.M., Chung E., Duncan M., Guzman E., Hartzell G., Hunnicke-Smith S., Hyman R., Kayser A., Komp C., Laskin-Kari D., Lew H., Lin D., Mosedale D., Nakahara K., Nathath A., Norgren R., Oefner P., Oh C., Petel F.X., Roberts D., Sehl P., Schramm S., Shoopen T., Smith V., Taylor P., Wei Y., Yeltos M., Botstein D., Davis R.W.; Submitted (DEC-1994) to the EMBL/GenBank/DDBJ databases.
 RN [3]
 RP CHARACTERIZATION, PubMed=9196079;
 RX MEDLINE=9733480;
 RA Russnak R., Pereira S., Platt T;
 RT "RNA binding analysis of yeast REF2 and its two-hybrid interaction with a new gene product, FIR1."
 RL Gene Expr. 6:24-258(1996);
 CC -!- FUNCTION: INTERACTS WITH POLY(A) POLYMERASE AND WITH REF2.

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CC EMBL: U17262; AAA46625; 1;
 DR EMBL: U18778; AAA64565; 1;
 DR SGG; S0000834; FIR1.
 FT CONFLICT 663 R -> P (IN REF 2);
 SQ SEQUENCE 925 AA; 104701 MW; 70798399EB31322B CRC64;

Query Match 80.5%; Score 33; DB 1; Length 925;
 Best Local Similarity 71.4%; Pred. No. 19;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 KKRMHC 7
 |||||:
 Db 841 KKRLSHC 847

RESULT 5
 ID FKBP_MOUSE STANDARD; PRT; 140 AA.
 AC P43678;
 DT 01-NOV-1995 (Rel. 32; Created)
 DT 01-JUL-1999 (Rel. 38; Last annotation update)
 DE FK506-BINDING PROTEIN PRECURSOR (FKBP-13) (PEPTIDYL-PROLYL CIS-TRANS ISOMERASE) (PPIASE) (EC 5.2.1.8);
 DE FKBP2 OR FKBP12.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 NCBI_TAXID=10909;
 OX [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=12/SVJ; TISSUE=Liver;
 RX MEDLINE=94085190; Pubmed=7505249;
 RA Herrickson B.A., Zhang W., Craig R.J., Jin Y.J., Bierer R.E., Burakoff S.J., Dilella A.G.;
 RA "Structural organization of the genes encoding human and murine FK506-binding protein (FKBP) 13 and comparison to FKBP1.";
 RT Gene 134:271-275(1993).
 CC -!- FUNCTION: PPIASE ACCELERATE THE FOLDING OF PROTEINS.
 CC -!- CATALYTIC ACTIVITY: CIS-TRANS ISOMERIZATION OF PROLINE IMIDIC PEPTIDE BONDS IN OLIGOPEPTIDES.
 CC -!- ENZYME REGULATION: INHIBITION BY BOTH FK506 AND RAPAMYCIN.
 CC -!- SUBCELLULAR LOCATION: ENDOPLASMIC RETICULUM LOREN. MEMBRANE ASSOCIATED (PROBABLE).
 CC -!- SIMILARITY: BELONGS TO THE FKBP-TYPE PPIASE FAMILY.
 CC DR M77831; AAA37631; 1;
 DR MGD: MGI:95542; FKBP2.
 DR InterPro: IPR001179; FKBP_PPIase.
 DR Pfam: PF0254; FKBP_PPIase.
 DR PROSITE: PS00453; FKBP_PPIASE_1; 1.
 DR PROSITE: PS00454; FKBP_PPIASE_2; 1.
 DR PROSITE; PS50059; FKBP_PPIASE_3; 1.
 DR Isomerase; Rotamase; Signal; Endoplasmic reticulum.
 FT SIGNAL_1 22

FT	CHAIN	23	140	FK506-BINDING PROTEIN.
FT	SITE	137	140	PREDICTED SECRETION FROM ER (POTENTIAL).
SEQ	SEQUENCE	140 AA;	15344 MW;	F4E7FCC7766A0416 CRC64;
Query Match	Score	32;	DB 1;	Length 140;
Best Local Similarity	78.0%	Score	32;	DB 1;
Matches	71.4%	Pred. No	4.8;	Length 140;
Matches	5;	Mismatches	1;	Indels
Qy	1 KKRIMHC 7	O;	Gaps	0;
Db	34 KKRVDH ₄ 40			
RESULT	6			
FKB2_HUMAN	STANDARD;	PRT;	141 AA.	
ID	FKB2_HUMAN	STANDARD;	PRT;	141 AA.
AC	P26885;			
DT	01-AUG-1992 (Rel. 23, Created)			
DT	01-AUG-1992 (Rel. 23, Last sequence update)			
DT	15-JUL-1999 (Rel. 38, Last annotation update)			
DE	FK506-BINDING PROTEIN PRECURSOR (FKBP-13) (PEPTIDYL-PROLYL CIS-TRANS ISOMERASE) (PPASE) (EC 5.2.1.8).			
GN	FKBP2 OR FKBP13.			
OS	Homo sapiens (Human).			
OU	Bukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.			
NCBI_TaxID	9606;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	TISSUE=Colon carcinoma;			
RA	LINEID=9131947; PubMed=1713687;			
JN	Y.-J. Albers M.W., Lane W.S., Bieser B.E., Schreiber S.L.,			
RA	Burakoff S.J.			
RT	"Molecular cloning of a membrane-associated human FK506- and rapamycin-binding protein, FKBP-13.";			
RP	PROC. NATL. ACAD. SCI. U.S.A. 88:6677-6681(1991).			
RN	[2]			
SEQUENCE FROM N.A.				
RP	LINEID=9131947; PubMed=1713687;			
RC	Dilella A.G., Hawkins A., Craig R.J., Schreiber S.L., Griffin C.A.;			
RA	"Chromosomal band assignments of the genes encoding human FKBP12 and FKBP13."			
RT	Biochem. Biophys. Res. Commun. 189:819-823(1992).			
CC	-1- FUNCTION: PPases ACCELERATE THE FOLDING OF PROTEINS.			
CC	-1- CATALYTIC ACTIVITY: CIS-TRANS ISOMERIZATION OF PROLINE IMIDIC PEPTIDE BONDS IN OLIGOPEPTIDES.			
CC	-1- ENZYME REGULATION: INHIBITED BY BOTH FK506 AND RAPAMYCIN.			
CC	-1- SUBCELLULAR LOCATION: ENDOPLASMIC RETICULUM LUMEN. MEMBRANE ASSOCIATED (PROBABLE).			
CC	-1- TISSUE SPECIFICITY: T-CELLS AND THYMUS.			
CC	-1- SIMILARITY: BELONGS TO THE FKBP-TYPE PPASE FAMILY.			
CC	-----			
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DR	EMBL: M65128; AAA5873.1;			
DR	EMBL: M75059; AAA3653.1;			
DR	PIR: JC1365; JC1365.			
DR	HSSP: P20001; IFKT.			
DR	MM: 186946;			
DR	InterPro: IPR001179; FKBP_PPASE.			
Pfam: PF00254; FKBP_1.				
DR	PROSITE: PS00454; FKBP_PPASE_1.			
DR	PROSITE: PS50059; FKBP_PPASE_2.			
DR	PROSITE: PS50059; FKBP_PPASE_3.			
DR	Isomerase; Rotamase.			
FT	1			
Qy	1 KKRIMHC 7			
Db	34 KKRVDH ₄ 40			
RESULT	6			
FKB2_HUMAN	STANDARD;	PRT;	141 AA.	
ID	FKB2_HUMAN	STANDARD;	PRT;	141 AA.
AC	P26885;			
DT	01-AUG-1992 (Rel. 23, Created)			
DT	01-AUG-1992 (Rel. 23, Last sequence update)			
DT	15-JUL-1999 (Rel. 38, Last annotation update)			
DE	FK506-BINDING PROTEIN PRECURSOR (FKBP-13) (PEPTIDYL-PROLYL CIS-TRANS ISOMERASE) (PPASE) (EC 5.2.1.8).			
GN	FKBP2 OR FKBP13.			
OS	Homo sapiens (Human).			
OU	Bukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.			
NCBI_TaxID	9606;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	SURAINK12 / TGL;			
RX	MEDLINE=9429472; PubMed=8021226;			
RA	Menon N.K., Chatelus C.Y., Dervartanian M., Wendt J.C., Bacteriaceae;			
RA	Escherichia; Proteobacteria; gamma subdivision; Enterobacteriaceae;			
RA	Escherichia.			
NCBI_TaxID	=562			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	SURAINK12 / TGL;			
RX	MEDLINE=9429472; PubMed=8021226;			
RA	Menon N.K., Chatelus C.Y., Dervartanian M., Wendt J.C., Bacteriaceae;			
RA	Escherichia; Proteobacteria; gamma subdivision; Enterobacteriaceae;			
RA	Escherichia.			
NCBI_TaxID	=562			
RN	[2]			
SEQUENCE FROM N.A.				
RP	LINEID=9131947; PubMed=1713687;			
RC	Jin Y.-J., Albers M.W., Lane W.S., Bieser B.E., Schreiber S.L.,			
RA	Burakoff S.J.			
RT	"Molecular cloning of a membrane-associated human FK506- and rapamycin-binding protein, FKBP-13.";			
RP	PROC. NATL. ACAD. SCI. U.S.A. 88:6677-6681(1991).			
RN	[2]			
SEQUENCE FROM N.A.				
RP	LINEID=9131947; PubMed=1713687;			
RC	Dilella A.G., Hawkins A., Craig R.J., Schreiber S.L., Griffin C.A.;			
RA	"Chromosomal band assignments of the genes encoding human FKBP12 and FKBP13."			
RT	Biochem. Biophys. Res. Commun. 189:819-823(1992).			
CC	-1- FUNCTION: PPases ACCELERATE THE FOLDING OF PROTEINS.			
CC	-1- CATALYTIC ACTIVITY: CIS-TRANS ISOMERIZATION OF PROLINE IMIDIC PEPTIDE BONDS IN OLIGOPEPTIDES.			
CC	-1- ENZYME REGULATION: INHIBITED BY BOTH FK506 AND RAPAMYCIN.			
CC	-1- SUBCELLULAR LOCATION: ENDOPLASMIC RETICULUM LUMEN. MEMBRANE ASSOCIATED (PROBABLE).			
CC	-1- TISSUE SPECIFICITY: T-CELLS AND THYMUS.			
CC	-1- SIMILARITY: BELONGS TO THE FKBP-TYPE PPASE FAMILY.			
CC	-----			
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DR	EMBL: M65128; AAA5873.1;			
DR	EMBL: M75059; AAA3653.1;			
DR	PIR: JC1365; JC1365.			
DR	HSSP: P20001; IFKT.			
DR	MM: 186946;			
DR	InterPro: IPR001179; FKBP_PPASE.			
Pfam: PF00254; FKBP_1.				
DR	PROSITE: PS00454; FKBP_PPASE_1.			
DR	PROSITE: PS50059; FKBP_PPASE_2.			
DR	PROSITE: PS50059; FKBP_PPASE_3.			
DR	Isomerase; Rotamase.			
FT	1			
Qy	1 KKRIMHC 7			
Db	34 KKRVDH ₄ 40			
RESULT	6			
FKB2_HUMAN	STANDARD;	PRT;	141 AA.	
ID	FKB2_HUMAN	STANDARD;	PRT;	141 AA.
AC	P26885;			
DT	01-AUG-1992 (Rel. 23, Created)			
DT	01-AUG-1992 (Rel. 23, Last sequence update)			
DT	15-JUL-1999 (Rel. 38, Last annotation update)			
DE	FK506-BINDING PROTEIN PRECURSOR (FKBP-13) (PEPTIDYL-PROLYL CIS-TRANS ISOMERASE) (PPASE) (EC 5.2.1.8).			
GN	FKBP2 OR FKBP13.			
OS	Homo sapiens (Human).			
OU	Bukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.			
NCBI_TaxID	9606;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	SURAINK12 / TGL;			
RX	MEDLINE=9429472; PubMed=8021226;			
RA	Menon N.K., Chatelus C.Y., Dervartanian M., Wendt J.C., Bacteriaceae;			
RA	Escherichia; Proteobacteria; gamma subdivision; Enterobacteriaceae;			
RA	Escherichia.			
NCBI_TaxID	=562			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	SURAINK12 / TGL;			
RX	MEDLINE=9429472; PubMed=8021226;			
RA	Menon N.K., Chatelus C.Y., Dervartanian M., Wendt J.C., Bacteriaceae;			
RA	Escherichia; Proteobacteria; gamma subdivision; Enterobacteriaceae;			
RA	Escherichia.			
NCBI_TaxID	=562			
RN	[2]			
SEQUENCE FROM N.A.				
RP	LINEID=9131947; PubMed=1713687;			
RC	Jin Y.-J., Albers M.W., Lane W.S., Bieser B.E., Schreiber S.L.,			
RA	Burakoff S.J.			
RT	"Molecular cloning of a membrane-associated human FK506- and rapamycin-binding protein, FKBP-13.";			
RP	PROC. NATL. ACAD. SCI. U.S.A. 88:6677-6681(1991).			
RN	[2]			
SEQUENCE FROM N.A.				
RP	LINEID=9131947; PubMed=1713687;			
RC	Dilella A.G., Hawkins A., Craig R.J., Schreiber S.L., Griffin C.A.;			
RA	"Chromosomal band assignments of the genes encoding human FKBP12 and FKBP13."			
RT	Biochem. Biophys. Res. Commun. 189:819-823(1992).			
CC	-1- FUNCTION: PPases ACCELERATE THE FOLDING OF PROTEINS.			
CC	-1- CATALYTIC ACTIVITY: CIS-TRANS ISOMERIZATION OF PROLINE IMIDIC PEPTIDE BONDS IN OLIGOPEPTIDES.			
CC	-1- ENZYME REGULATION: INHIBITED BY BOTH FK506 AND RAPAMYCIN.			
CC	-1- SUBCELLULAR LOCATION: ENDOPLASMIC RETICULUM LUMEN. MEMBRANE ASSOCIATED (PROBABLE).			
CC	-1- TISSUE SPECIFICITY: T-CELLS AND THYMUS.			
CC	-1- SIMILARITY: BELONGS TO THE FKBP-TYPE PPASE FAMILY.			
CC	-----			
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DR	EMBL: M65128; AAA5873.1;			
DR	EMBL: M75059; AAA3653.1;			
DR	PIR: JC1365; JC1365.			
DR	HSSP: P20001; IFKT.			
DR	MM: 186946;			
DR	InterPro: IPR001179; FKBP_PPASE.			
Pfam: PF00254; FKBP_1.				
DR	PROSITE: PS00454; FKBP_PPASE_1.			
DR	PROSITE: PS50059; FKBP_PPASE_2.			
DR	PROSITE: PS50059; FKBP_PPASE_3.			
DR	Isomerase; Rotamase.			
FT	1			
Qy	1 KKRIMHC 7			
Db	34 KKRVDH ₄ 40			
RESULT	6			
FKB2_HUMAN	STANDARD;	PRT;	141 AA.	
ID	FKB2_HUMAN	STANDARD;	PRT;	141 AA.
AC	P26885;			
DT	01-AUG-1992 (Rel. 23, Created)			
DT	01-AUG-1992 (Rel. 23, Last sequence update)			
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GN	FKBP2 OR FKBP13.			
OS	Homo sapiens (Human).			
OU	Bukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.			
NCBI_TaxID	9606;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	SURAINK12 / TGL;			
RX	MEDLINE=9429472; PubMed=8021226;			
RA	Menon N.K., Chatelus C.Y., Dervartanian M., Wendt J.C., Bacteriaceae;			
RA	Escherichia; Proteobacteria; gamma subdivision; Enterobacteriaceae;			
RA	Escherichia.			
NCBI_TaxID	=562			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	SURAINK12 / TGL;			
RX	MEDLINE=9429472; PubMed=8021226;			
RA	Menon N.K., Chatelus C.Y., Dervartanian M., Wendt J.C., Bacteriaceae;			
RA	Escherichia; Proteobacteria; gamma subdivision; Enterobacteriaceae;			
RA	Escherichia.			
NCBI_TaxID	=562			
RN	[2]			
SEQUENCE FROM N.A.				
RP	LINEID=9131947; PubMed=1713687;			
RC	Jin Y.-J., Albers M.W., Lane W.S., Bieser B.E., Schreiber S.L.,			
RA	Burakoff S.J.			
RT	"Molecular cloning of a membrane-associated human FK506- and rapamycin-binding protein, FKBP-13.";			
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RN	[2]			
SEQUENCE FROM N.A.				
RP	LINEID=9131947; PubMed=1713687;			
RC	Dilella A.G., Hawkins A., Craig R.J., Schreiber S.L., Griffin C.A.;			
RA	"Chromosomal band assignments of the genes encoding human FKBP12 and FKBP13."			
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CC	-1- ENZYME REGULATION: INHIBITED BY BOTH FK506 AND RAPAMYCIN.			
CC	-1- SUBCELLULAR LOCATION: ENDOPLASMIC RETICULUM LUMEN. MEMBRANE ASSOCIATED (PROBABLE).			
CC	-1- TISSUE SPECIFICITY: T-CELLS AND THYMUS.			
CC	-1- SIMILARITY: BELONGS TO THE FKBP-TYPE PPASE FAMILY.			
CC	-----			
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DR	EMBL: M65128; AAA5873.1;			
DR	EMBL: M75059; AAA3653.1;			
DR	PIR: JC1365; JC1365.			
DR	HSSP: P20001; IFKT.			
DR	MM: 186946;			
DR	InterPro: IPR001179; FKBP_PPASE.			
Pfam: PF00254; FKBP_1.				
DR	PROSITE: PS00454; FKBP_PPASE_1.			
DR	PROSITE: PS50059; FKBP_PPASE_2.			
DR	PROSITE: PS50059; FKBP_PPASE_3.			
DR	Isomerase; Rotamase.			
FT	1			
Qy	1 KKRIMHC 7			
Db	34 KKRVDH ₄ 40			
RESULT	6			
FKB2_HUMAN	STANDARD;	PRT;	141 AA.	
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DT	01-AUG-1992 (Rel. 23, Last sequence update)			
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GN	FKBP2 OR FKBP13.			
OS	Homo sapiens (Human).			
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OC	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.			
NCBI_TaxID	9606;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	SURAINK12 / TGL;			
RX	MEDLINE=9429472; PubMed=8021226;			
RA	Menon N.K., Chatelus C.Y., Dervartanian M., Wendt J.C., Bacteriaceae;			
RA	Escherichia; Proteobacteria; gamma subdivision; Enterobacteriaceae;			
RA	Escherichia.			
NCBI_TaxID	=562			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	SURAINK12 / TGL;			
RX	MEDLINE=9429472; PubMed=8021226;			
RA	Menon N.K., Chatelus C.Y., Dervartanian M., Wendt J.C., Bacteriaceae;			
RA	Escherichia; Proteobacteria; gamma subdivision; Enterobacteriaceae;			
RA	Escherichia.			
NCBI_TaxID	=562			
RN	[2]			
SEQUENCE FROM N.A.				
RP	LINEID=9131947; PubMed=1713687;			
RC	Jin Y.-J., Albers M.W., Lane W.S., Bieser B.E., Schreiber S.L.,			
RA	Burakoff S.J.		</	

"Complete genome sequence of enterohemorrhagic Escherichia coli O157:H7 and genomic comparison with a laboratory strain K-12.";

RT DNA Res. 8:11-22(2001).

RL -!- FUNCTION: PARTICIPATES IN THE PERIPLASMIC ELECTRON-TRANSFERRING ACTIVITY OF HYDROGENASE 2 DURING ITS CATALYTIC TURNOVER.

CC -!- COFACTOR: BINDS 3 4Fe-4S CLUSTERS AND A 3Fe-4S CLUSTER.

CC -!- SUBCELLULAR LOCATION: PERIPLASMIC.

CC -!- SIMILARITY: THE IRON-SULFUR CENTERS ARE SIMILAR TO THOSE OF

CC 'BACTERIAL-TYPE' 4Fe-4S FERREDOXINS. CC -!- CAUTION: WAS ORIGINALLY (REF.1) THOUGHT TO BE THE SMALL SUBUNIT OF HYDROGENASE 2.

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CC -!- HSPSP: P59071; 1XER.

DR EcoGene: EG11799; hybA.

DR InterPro: IPR01450; 4Fe4S_ferritin.

DR Pfam: PF00037; fer4; 1.

DR PROSITE: PS00198; 4FEIS_FERREDOXIN; 1.

DR Oxidoreductase; Signal; Periplasmic; Iron-sulfur; 4Fe-4S; 3Fe-4S;

KW Complete proteome.

FT SIGNAL 1 27 POTENTIAL.

FT CHAIN 28 328 HYDROGENASE-2 OPERON PROTEIN_HBA.

FT METAL 47 47 IRON-SULFUR 1 (4Fe-4S) (POTENTIAL).

FT METAL 50 50 IRON-SULFUR 1 (4Fe-4S) (POTENTIAL).

FT METAL 53 53 IRON-SULFUR 1 (4Fe-4S) (POTENTIAL).

FT METAL 57 57 IRON-SULFUR 1 (4Fe-4S) (POTENTIAL).

FT METAL 112 112 IRON-SULFUR 2 (3Fe-4S) (POTENTIAL).

FT METAL 115 115 IRON-SULFUR 2 (3Fe-4S) (POTENTIAL).

FT METAL 120 120 IRON-SULFUR 2 (3Fe-4S) (POTENTIAL).

FT METAL 124 124 IRON-SULFUR 2 (3Fe-4S) (POTENTIAL).

FT METAL 145 145 IRON-SULFUR 3 (4Fe-4S) (POTENTIAL).

FT METAL 148 148 IRON-SULFUR 3 (4Fe-4S) (POTENTIAL).

FT METAL 151 151 IRON-SULFUR 3 (4Fe-4S) (POTENTIAL).

FT METAL 155 155 IRON-SULFUR 3 (4Fe-4S) (POTENTIAL).

FT METAL 174 174 IRON-SULFUR 4 (4Fe-4S) (POTENTIAL).

FT METAL 177 177 IRON-SULFUR 4 (4Fe-4S) (POTENTIAL).

FT METAL 193 193 IRON-SULFUR 4 (4Fe-4S) (POTENTIAL).

FT METAL 197 197 IRON-SULFUR 4 (4Fe-4S) (POTENTIAL).

SQ SEQUENCE 328 AA; 36003 MW; 77203AF05061662 CRC64;

Query Match Score 32; DB 1; Length 328;

Best Local Similarity 71.4%; Pred. No. 11; Mismatches 1; Indels 0; Gaps 0;

Qy 1 KKrimhc 7

Db 109 KKQCMHC 115

RESULT 8
DYR2_HUMAN STANDARD; PRT: 528 AA.
ID Q92630;
AC DT 20-AUG-2001 (Rel. 40, Created)
DT 20-AUG-2001 (Rel. 40, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE DUAL-SPECIFICITY TYROSINE-PHOSPHORYLATION REGULATED KINASE 2
DE (EC 2.7.1.-).
DE DYR2.
GN Homo sapiens (Human).

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
OC NCBI_TAXID=9606;
OX [1]
RN SEQUENCE FROM N.A.
RP TISSUE=Brian;
RC MEDLINE=98421512; PubMed=9748265;
RX RA Becker W., Weber Y., Wetzel R., Eirrmber K., Tejedor F.J.,
RA Joost H.-G.;
RN "Sequence characteristics, subcellular localization, and substrate specificity of DYRK related kinases,";
RT specificity protein kinases.;
RT "Sequence characteristics, subcellular localization, and substrate specificity of DYRK related kinases,";
RT sequence of dual
SEQUENCE OF 320-528 FROM N.A.
RP TISSUE=Placenta;
RA Becker W., Joost H.-G.;
RL Submitted (NOV-1996) to the EMBL/GenBank/DDBJ databases.
CC -!- FUNCTION: IN VITRO; CAN PHOSPHORYLATE HISTONES H3 AND H2B ON SER AND THR RESIDUES. MAY BE INVOLVED IN THE REGULATION OF CELLULAR GROWTH AND/OR DEVELOPMENT.
CC -!- SUBCELLULAR LOCATION: CYTOPLASMIC.
CC -!- PTM: AUTOPHOSPHORYLATION ON TYR RESIDUES.
CC -!- SIMILARITY: BELONGS TO THE SER/THR FAMILY OF PROTEIN KINASES.
CC MNK/DYRK SUBFAMILY.
CC
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CC
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CC
CC DR EMBL: Y13493; CAA73985; 1;
CC DR EMBL: Y09216; CAA70418; 1;
CC DR HSSP: Q165539; 1WFC.
CC DR NMIM: 603496; -;
CC DR InterPro: IPR000719; Euk_Pkinase.
CC DR InterPro: IPR000729; Ser_Thr_kin_actsite.
CC DR Pfam: PF00569; Pkinase; 2.
CC DR SMART: SMM00220; S-TKc; 1.
CC DR PROSITE: PS00107; PROTEIN_KINASE_ATP; 1.
CC DR PROSITE: PS55011; PROTEIN_KINASE_DOM; 1.
CC DR PROSITE: PS00108; PROTEIN_KINASE_ST; 1.
CC DR Transferase: Serine/threonine-protein kinase; Tyrosine-protein kinase; ATP-binding; Phosphorylation.
CC KW DOMAIN 149 462 PROTEIN KINASE.
CC FT BINDING 155 163 ATP (BY SIMILARITY).
CC FT BINDING 178 178 ATP (BY SIMILARITY).
CC FT ACT SITE 275 275 BY SIMILARITY.
CC SQ SEQUENCE 528 AA; 59714 MW; AF26822DD9522D7 CRC64;

Query Match Score 32; DB 1; Length 528;
Best Local Similarity 71.4%; Pred. No. 18; Mismatches 5; Conservative 1; Indels 0; Gaps 0;

Qy 1 KKrimhc 7

Db 268 KNRIHC 274

RESULT 9
AMP1_LYCES STANDARD; PRT: 571 AA.
ID Q10712; Q959A3;
AC DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE CHLOROPLAST AMINOPEPTIDASE 1 PRECURSOR (EC 3.4.11.1) (LEUCINE AMINOPEPTIDASE) (PROLINE AMINOPEPTIDASE) (PR57).
DE DE AMINOPEPTIDASE (LAP) (LEUCYL AMINOPEPTIDASE) (PROLINE AMINOPEPTIDASE) (PR57).

GN LAPA1 OR LAP OR LAP2.
 OS Lycopersicon esculentum (Tomato).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; eudicots; core eudicots;
 OC Asteridae; euasterids I; Solanales; Solanaceae; Solanum.
 OX NCBI_TAXID=4081;
 RN [1] RP SEQUENCE FROM N.A.
 RC STRAIN=CV; PETO 238R; TISSUE=Leaf;
 RX MEDLINE=9641572; PubMed=88202420;
 RA Gu Y.Q., Chao W.S., Walling L.L.;
 RT "Localization and post-translational processing of the wound-induced
 leucine aminopeptidase proteins of tomato.";
 RL J. Biol. Chem. 271:25880-25887(1996).
 RN [2] RP SEQUENCE OF 49-571 FROM N.A.
 RC STRAIN=CV; VF36; TISSUE=Pistil;
 RX MEDLINE=94052201; PubMed=8334314;
 RA Pautz V., Holzer F.M., Reisch B., Walling L.L.;
 RT "Leucine aminopeptidase: an inducible component of the defense
 response in Lycopersicon esculentum (tomato)." ;
 RL Proc. Natl. Acad. Sci. U. S. A. 90:9906-9910(1993).
 CC -1- FUNCTION: PRESUMABLY INVOLVED IN THE PROCESSING AND REGULAR
 CC -1- TURNOVER OF INTRACELLULAR PROTEINS.
 CC -1- CATALYTIC ACTIVITY: RELEASE OF AN N-TERMINAL AMINO ACID, XAA-|-
 CC XBB-, IN WHICH XAA IS PREFERABLY LEU, BUT MAY BE OTHER AMINO ACIDS
 CC -1- INCLUDING PRO ALTHOUGH NOT ARG OR LYS, AND XBB MAY BE PRO.
 CC -1- COFACTOR: BINDS TWO ZINC IONS (BY SIMILARITY).
 CC -1- INDUCTION: BY WOUNDING.
 CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY M17: ALSO KNOWN AS THE
 CC CYTOSOL AMINOPETIDASE FAMILY.
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 CC or send an email to license@isb-sib.ch).
 DR U50151; AAC49456; 1;
 EMBL; U5052; AAC19457; 1;
 DR U20593; AAC80498; 1;
 DR HSSP; P00727; 1BPM.
 DR MEROPS; M17.002;
 DR InterPro; IP000819; Peptidase_M17; 1.
 DR PRANTS; PRO0481; LAMNOPPTDSE.
 DR PROSITE; PS00631; CYTOSOLAP; 1.
 KW TRANSIT Peptide: Chloroplast; Aminopeptidase; Hydrolase; Zinc.
 FT TRANSIT 1 53 CHLOROPLAST (POTENTIAL).
 FT CHAIN 54 571 CHLOROPLAST AMINOPETIDASE 1.
 FT DOMAIN 169 174 POLY-ALA.
 FT METAL 342 342 ZINC (2) (BY SIMILARITY).
 FT METAL 347 347 ZINC (1 AND 2) (BY SIMILARITY).
 FT METAL 367 367 ZINC (2) (BY SIMILARITY).
 FT METAL 427 427 ZINC (1) (BY SIMILARITY).
 FT METAL 429 429 ZINC (1 AND 2) (BY SIMILARITY).
 FT ACT-SITE 354 354 POTENTIAL.
 FT VARIANT 358 358 R -> G.
 FT CONFLICT 271 271 P -> S (IN REF. 2).
 FT CONFLICT 315 315 T -> S (IN REF. 3).
 FT CONFLICT 515 515 T -> L (IN CLEONE PBLAP2).
 SQ 571 AA; 60279 MW; C7A224837E/3939D CRC64;

Query Match 78.0%; Score 32; DB 1; Length 571;
 Best Local Similarity 83.3%; Pred. No. 20;
 Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 KRIMHC 7
 Dib 48 KRVIC 53

RESULT 10
 ID AMPL_SOLTU STANDARD;
 ID AMPL_SOLTU PRT;
 AC P3127;
 AC P3127;
 DT 01-JUL-1993 (Rel. 26, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 01-OCT-1996 (Rel. 34, Last annotation update)
 DE CHLOROPLAST AMINOPETIDASE PRECURSOR (EC 3.4.11.1) (LEUCINE
 DE AMINOPETIDASE) (LAP) (LEUCYL AMINOPETIDASE) (PROLINE AMINOPETIDASE)
 DE (EC 3.4.11.5) (PROLYL AMINOPETIDASE)
 DE LAP
 GN Solanum tuberosum (Potato).
 OS Solanaceae; Solanum tuberosum; Solanaceae; Solanum.
 OC Eukaryote; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; eudicots; core eudicots;
 OC Asteridae; euasterids I; Solanales; Solanaceae; Solanum.
 OC NCBI_TAXID=4113;
 RN [1] RP SEQUENCE FROM N.A.
 RC STRAIN=CV DESTRIE; TISSUE=leaf;
 RX MEDLINE=94339796; PubMed=7765119;
 RA Herbers K., Prat S., Willmitzer L.;
 RT "Functional analysis of a leucine aminopeptidase from Solanum
 tuberosum L." ;
 RL Planta 194:230-240(1994).
 RN [2] RP SEQUENCE OF 19-573 FROM N.A.
 RC STRAIN=CV DESTRIE; TISSUE=leaf;
 RX MEDLINE=94005746; PubMed=1392612;
 RA Hildmann T., Ebnet M., Pena-Cortes H., Sanchez-Serrano J.J.,
 RA Willmitzer L., Prat S.;
 RT "General roles of abscisic and jasmonic acids in gene activation as a
 result of mechanical wounding." ;
 RL Plant Cell 4:1157-1170(1992).
 CC -1- FUNCTION: PRESUMABLY INVOLVED IN THE PROCESSING AND REGULAR
 CC -1- TURNOVER OF INTRACELLULAR PROTEINS.
 CC -1- CATALYTIC ACTIVITY: RELEASE OF AN N-TERMINAL AMINO ACID, XAA-|-
 CC XBB-, IN WHICH XAA IS PREFERABLY LEU, BUT MAY BE OTHER AMINO ACIDS
 CC -1- INCLUDING PRO ALTHOUGH NOT ARG OR LYS, AND XBB MAY BE PRO.
 CC -1- COFACTOR: BINDS TWO ZINC IONS (BY SIMILARITY).
 CC -1- SUBCELLULAR LOCATION: CHLOROPLAST (PROBABLE).
 CC -1- SUBCELLULAR LOCATION: CHLOROPLAST (BY SIMILARITY).
 CC -1- TISSUE SPECIFICITY: IN TUBERS AND FLORAL BUDS OF UNTRIMMED PLANTS.
 CC -1- COFACTOR: BINDS TWO ZINC IONS (BY SIMILARITY).
 CC -1- SUBUNIT: HOMOHEXAMER (PROBABLE).
 CC -1- SUBUNIT: HOMOHEXAMER (PROBABLE).
 CC -1- TISSUE SPECIFICITY: IN TUBERS AND FLORAL BUDS OF UNTRIMMED PLANTS.
 CC -1- AFTERABA TREATMENT OR MECHANICAL WOUNDING IS MOSTLY ACCUMULATED
 CC -1- IN LEAVES, TO A LESSER EXTENT IN STEMS, BUT NOT IN ROOTS.
 CC -1- INDUCTION: BY ABSICNIC ACID (ABA), JASMONIC ACID (JA) AND
 CC WOUNDING.
 CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY M17; ALSO KNOWN AS THE
 CC CYTOSOL AMINOPETIDASE FAMILY.

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 or send an email to license@isb-sib.ch).

EMBL; Y77015; CAA54314.1;
 EMBL; X67645; CAA48038.1;
 DR PIR; S24769; S24769;
 DR PIR; PQ0470; PQ0470;
 DR HSSP; P00727; 1LAN.

DR	MEROPS; M17_002;	Qy	1 KRIMHC 7
DR	InterPro; IPR000819; Peptidase_M17;	Pfam; PF00883; Peptidase_M17;	1 : 1
DR	PRINTS; PRO048I; LAMNOPEPTIDASE;	Db	5 KKMKMH C 11
DR	PROSITE; PS00631; CYTOSOL_AP; 1.		
KW	Transit peptidase; Chloroplast; Aminopeptidase; Hydrolase; Zinc;		
FT	TRANSIT 1	RESULT 12	
FT	CHAIN 53	Y091_NPYOP	STANDARD;
FT	CHLOROPLAST (POTENTIAL)	ID Y091_NPYOP	PRT;
FT	CHLOROPLAST AMINOPEPTIDASE.	010341;	279 AA.
FT	POLY-ALA.	AC DT	
DOMAIN	169 174	01 NOV-1997 (Rel. 35, Created)	
FT	METAL 342	DT 01 NOV-1997 (Rel. 35, Last sequence update)	
FT	ZINC (2) (BY SIMILARITY).	DT 01 NOV-1997 (Rel. 35, Last annotation update)	
FT	METAL 347	HYPOTHETICAL 29.3 KDa PROTEIN (ORF92).	
FT	ZINC (1 AND 2) (BY SIMILARITY).	DE OS	Orgyia pseudotsugata multicapsid polyhedrosis virus (OpMNPV).
FT	METAL 367	OC Viruses; dsDNA viruses, no RNA stage; Baculoviridae;	
FT	ZINC (2) (BY SIMILARITY).	OC Nucleopolyhedrovirus.	
FT	METAL 427	OC NCBI_TaxID=164623;	
FT	ZINC (1) (BY SIMILARITY).	OX	
FT	METAL 429	RN [1]	
FT	ZINC (1 AND 2) (BY SIMILARITY).	RP SEQUENCE FROM N.A.	
ACT_SITE	POTENTIAL.	MEDLINE:9721300; PubMed:9126251;	
FT	354	RX	
ACT_SITE	431	RA Ahrens C.H., Russell R.R., Funk C.J., Evans J., Harwood S.,	
SEQUENCE	431 AA;	RA Rohrmann G.F.;	
SEQUENCE	573 AA;	RT "The sequence of the Orgyia pseudotsugata multicapsid nuclear	
	60122 MW;	RT polyhedrosis virus genome. ;	
	3152145A4A7FB291 CRC64;	RL Virology 229:381-399(1997).	
		CC -1- SIMILARITY: TO CORRESPONDING ORF IN ACMPNPV.	
		CC CC	
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		use by non-profit institutions as long as its content is in no way	
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		entities requires a license agreement (See http://www.isb-sib.ch/announce/	
		or send an email to license@isb-sib.ch).	
		CC CC	
		EMBL: U75930; AAC59091.1; -	
		DR Hypothetical protein.	
		KW SEQUENCE 279 AA; 29289 MW; 6FA4DA01009DBFO CRC64;	
		CC CC	
		Query Match 78.0%; Score 32; DB 1; Length 573;	
Best Local Similarity	83.3%;	Pred. No. 20;	
Matches 5;	Conservative 1;	Mismatches 0;	
		Indels 0;	
		Gaps 0;	
Qy	2 KRIMHC 7		
Db	48 KRIVHC 53		
		RESULT 11	
		RL44_TRYBB	STANDARD;
		ID RL44_TRYBB	PRT;
		AC P17813;	105 AA.
		DT 01-AUG-1990 (Rel. 15, Created)	
		DT 20-AUG-2001 (Rel. 40, Last sequence update)	
		DT 20-AUG-2001 (Rel. 40, Last annotation update)	
		DE 60S RIBOSOMAL PROTEIN L44.	
		GN RPL44.	
		OS Trypanosoma brucei brucei.	
		OC Euglenozoa; Kinetoplastida; Trypanosomatidae; Trypanosoma.	
		NCBI_TaxID=5702;	
		RN [1]	
		RP SEQUENCE FROM N.A.	
		RC STRAIN=EATRO 1125;	
		RX MEDLINE=90251460; PubMed=2339065;	
		RA Tebabi P., Halleux S., Pays E.,	
		RT Nucleotide sequence of a full-length cDNA coding for the ribosomal L44 protein of Trypanosoma brucei ¹ .	
		RL Nucleic Acids Res. 18:2809-2819 (1990).	
		CC -1- SUBCELLULAR LOCATION: CYTOPLASMIC.	
		CC -1- SIMILARITY: BELONGS TO THE L44E FAMILY OF RIBOSOMAL PROTEINS.	
		CC CC	
		This SWISS-PROT entry is copyright. It is produced through a collaboration	
		between the Swiss Institute of Bioinformatics and the EMBL outstation -	
		the European Bioinformatics Institute. There are no restrictions on its	
		use by non-profit institutions as long as its content is in no way	
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		entities requires a license agreement (See http://www.isb-sib.ch/announce/	
		or send an email to license@isb-sib.ch).	
		CC CC	
		EMBL: X52122; CAA36167.1; -	
		DR PIR; S10012; R0UT6A.	
		DR InterPro; IPR000552; Ribosomal_L44_E.	
		DR Pfam; PF00935; Ribosomal_L44_2.	
		DR ProDom; PD002841; Ribosomal_L44E_1.	
		KW Ribosomal protein.	
		FT INIT_MET 0 0 BY SIMILARITY.	
		SEQUENCE 105 AA; 12322 MW; FA9423F109E7819 CRC64;	
		CC CC	
		Query Match 75.6%; Score 31; DB 1; Length 105;	
Best Local Similarity	71.4%;	Pred. No. 5.9;	
Matches 5;	Conservative 1;	Mismatches 1;	
		Indels 0;	
		Gaps 0;	

"P40K015 associates with p36 subunit and requires both nuclear translocation and Thr176 phosphorylation to generate cdk-activating kinase activity in Xenopus oocytes.";

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RC TISSUE=Brain;
 RA Yu W.; Sarginson J.; Gibbs R.A.; Submitted (MAR 1998) to the EMBL/GenBank/DDBJ databases.
 RL -!
 CC -! SIMILARITY: BELONGS TO THE CULLIN FAMILY.
 CC
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 CC or send an email to license@isb-sib.ch).
 CC
 DR EMBL; AF064087; AAC36304_1;
 DR EMBL; AB014517; BAA31592_1;
 DR EMBL; AF002537; AAC36682_1;
 DR EMBL; U58089; AAC50546_1;
 DR EMBL; AF050147; AAC28621_1;
 DR MIM; 603116;
 DR InterPro; IPR001373; Cullin;
 DR InterPro; IPR002119; Histone_H2A;
 DR Pfam; PF00888; Cullin; 1.
 DR ProDom; PDO00505; Histone_H2A; 1.
 DR SMART; SM00182; CULLIN; 1.
 DR PROSITE; PS01236; CULLIN_1; 1.
 DR PROSITE; PS50063; CULLIN_2; 1.
 FT CONFLICT 13 D -> G (IN REF. 3)
 FT CONFLICT 159 179 DHLRQILLNTRERKGEVVD -> GSSTANSTGQYDCRAE
 FT CONFLICT 426 451 RRSSRS (IN REF. 3).
 FT CONFLICT 768 AA; 88930 MW; DVFERVYKOHIAARRLLTKNSVSDSE -> MYLNVIINNTW
 SQ SEQUENCE QGDFSQSIKVILMLK (IN REF. 5).
 CRC64;

Query Match 75.6%; Score 31; DB 1; Length 768;
 Best Local Similarity 66.7%; Pred. No. 43; Mismatches 0; Indels 0; Gaps 0;
 Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
 Qy 2 KRMHC 7
 Db 246 ERVMHC 251

Search completed: February 12, 2002, 12:04:02
 Job time: 798 sec

Copyright (c) 1993 - 2000 CompuGen Ltd.	GenCore version 4.5	30	31	75.6	383	2	S51651
OM protein - protein search, using sw model		31	31	75.6	451	1	F64155
Run on:	February 12, 2002, 11:51:39 ; Search time 55.4 Seconds (without alignments) 9.625 Million cell updates/sec	32	31	75.6	476	2	T17330
Title:	US-09-606-129A-18	33	31	75.6	504	2	JG775
Perfect score:	41	34	31	75.6	504	2	A56534
Sequence:	1 KKRMHC 7	35	31	75.6	861	2	T41945
Scoring table:	BLOSUM62 Gapped 10.0 , Gapext 0.5	36	31	75.6	905	2	B21562
Searched:	219241 seqs, 76174552 residues	37	31	75.6	905	2	H87174
Total number of hits satisfying chosen parameters:	219241	38	31	75.6	1687	2	T0176
Minimum DB seq length: 0		39	30	73.2	108	2	EGF repeat transme
Maximum DB seq length: 2000000000		40	30	73.2	108	2	protein-tyrosine-p
Post-processing: Minimum Match 0%		41	30	73.2	269	2	hypothetical prote
Maximum Match 100%		42	30	73.2	370	2	conserved hypot
Listing first 45 summaries		43	30	73.2	457	2	probable oxoglutar
Database :	PIR-68.* 1: pir1: 2: pir2: 3: pir3: 4: pir4: 5: pir5: 6: pir6: 7: pir7: 8: pir8: 9: pir9: 10: pir10: 11: pir11: 12: pir12: 13: pir13: 14: pir14: 15: pir15: 16: pir16: 17: pir17: 18: pir18: 19: pir19: 20: pir20: 21: pir21: 22: pir22: 23: pir23: 24: pir24: 25: pir25: 26: pir26: 27: pir27: 28: pir28: 29: pir29:	44	30	73.2	508	2	2-oxoglutamate de
Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being Printed, and is derived by analysis of the total score distribution.		45	30	73.2	522	2	EGF repeat transme
							CDS protein F9L11.
ALIGNMENTS							
RESULT	1						
A42268		biliverdin reductase (EC 1.3.1.24) - rat					
C:Species: Rattus norvegicus (Norway rat)							
C:Date: 04-Mar-1993 #sequence_revision 18-Nov-1994 #text_change 05-Nov-1999							
C:Accession: A42268							
R:Fakhrai, H.; Maines, M.D.							
J. Biol. Chem. 267, 4023-4029, 1992							
A:Title: Expression and characterization of a cDNA for rat kidney biliverdin reduc							
A:Reference number: A42268; MUID:92156147							
A:Status: preliminary; not compared with conceptual translation							
A:Molecule type: nucleic acid; protein							
A:Residues: 1-295 <Pfam>							
A:Cross references: GB:MB1681; NID:9203177; PID:AAA40830.1; PID:g203178							
A:Experimental source: kidney							
A:Note: sequence extracted from NCBI backbone (NCBIP:82800)							
C:Keywords: liver; oxidoreductase							
RESULT	2						
Query Match Similarity 100.0%; Score 41; DB 2; Length 295;							
Best Local Similarity 100.0%; Pred. No. 0.36%; Mismatches 0; Indels 0; Gaps 0;							
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;							
Qy 1 KKRMHC 7							
Db 274 KKRMHC 280							
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Query Match Similarity 100.0%; Score 41; DB 2; Length 295;							
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Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;							
Qy 1 KKRMHC 7							
Db 274 KKRMHC 280							
RESULT	2						
Query Match Similarity 100.0%; Score 41; DB 2; Length 295;							
Best Local Similarity 100.0%; Pred. No. 0.36%; Mismatches 0; Indels 0; Gaps 0;							
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;							
Qy 1 KKRMHC 7							
Db 274 KKRMHC 280							
RESULT	2						
Query Match Similarity 100.0%; Score 41; DB 2; Length 295;							
Best Local Similarity 100.0%; Pred. No. 0.73%; Mismatches 0; Indels 0; Gaps 0;							
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;							
Qy 1 KKRMHC 7							
Db 274 KKRMHC 280							
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Query Match Similarity 100.0%; Score 41; DB 2; Length 295;							
Best Local Similarity 100.0%; Pred. No. 0.73%; Mismatches 0; Indels 0; Gaps 0;							
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;							
Qy 1 KKRMHC 7							
Db 274 KKRMHC 280							
RESULT	2						
Query Match Similarity 100.0%; Score 41; DB 2; Length 295;							
Best Local Similarity 100.0%; Pred. No. 0.73%; Mismatches 0; Indels 0; Gaps 0;							
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;							
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Db 274 KKRMHC 280							
RESULT	2						
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Best Local Similarity 100.0%; Pred. No. 0.73%; Mismatches 0; Indels 0; Gaps 0;							
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Qy 1 KKRMHC 7							
Db 274 KKRMHC 280							
RESULT	2						
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Best Local Similarity 100.0%; Pred. No. 0.73%; Mismatches 0; Indels 0; Gaps 0;							
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;							
Qy 1 KKRMHC 7							
Db 274 KKRMHC 280							
RESULT	2						
Query Match Similarity 100.0%; Score 41; DB 2; Length 295;							
Best Local Similarity 100.0%; Pred. No. 0.73%; Mismatches 0; Indels 0; Gaps 0;							
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;							
Qy 1 KKRMHC 7							
Db 274 KKRMHC 280							
RESULT	2						
Query Match Similarity 100.0%; Score 41; DB 2; Length 295;							
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Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;							
Qy 1 KKRMHC 7							
Db 274 KKRMHC 280							
RESULT	2						
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Db 274 KKRMHC 280							
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Best Local Similarity 100.0%; Pred. No. 0.73%; Mismatches 0; Indels 0; Gaps 0;							
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Qy 1 KKRMHC 7							
Db 274 KKRMHC 280							
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Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;							
Qy 1 KKRMHC 7							
Db 274 KKRMHC 280							
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Best Local Similarity 100.0%; Pred. No. 0.73%; Mismatches 0; Indels 0; Gaps 0;							
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;							
Qy 1 KKRMHC 7							
Db 274 KKRMHC 280							
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Query Match Similarity 100.0%; Score 41; DB 2; Length 295;							
Best Local Similarity 100.0%; Pred. No. 0.73%; Mismatches 0; Indels 0; Gaps 0;							
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;							
Qy 1 KKRMHC 7							
Db 274 KKRMHC 280							
RESULT	2						
Query Match Similarity 100.0%; Score 41; DB 2; Length 295;							
Best Local Similarity 100.0%; Pred. No. 0.73%; Mismatches 0; Indels 0; Gaps 0;							
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;							
Qy 1 KKRMHC 7							
Db 274 KKRMHC 280							
RESULT	2						
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Best Local Similarity 100.0%; Pred. No. 0.73%; Mismatches 0; Indels 0; Gaps 0;							
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Qy 1 KKRMHC 7							
Db 274 KKRMHC 280							
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Best Local Similarity 100.0%; Pred. No. 0.73%; Mismatches 0; Indels 0; Gaps 0;							
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;							
Qy 1 KKRMHC 7							
Db 274 KKRMHC 280							
RESULT	2						
Query Match Similarity 100.0%; Score 41; DB 2; Length 295;							
Best Local Similarity 100.0%; Pred. No. 0.73%; Mismatches 0; Indels 0; Gaps 0;							
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;							
Qy 1 KKRMHC 7							
Db 274 KKRMHC 280							
RESULT	2						
Query Match Similarity 100.0%; Score 41; DB 2; Length 295;							
Best Local Similarity 100.0%; Pred. No. 0.73%; Mismatches 0; Indels 0; Gaps 0;							
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;							
Qy 1 KKRMHC 7							
Db 274 KKRMHC 280							
RESULT	2						
Query Match Similarity 100.0%; Score 41; DB 2; Length 295;							
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Db 274 KKRMHC 280							
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Db 274 KKRMHC 280							
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Db 274 KKRMHC 280							
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Qy 1 KKRMHC 7							
Db 274 KKRMHC 280							
RESULT	2						
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Best Local Similarity 100.0%; Pred. No. 0.73%; Mismatches 0; Indels 0; Gaps 0;							
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;							
Qy 1 KKRMHC 7</							

RESULT 5
H86206 hypothetical protein [imported] - Arabidopsis thaliana
C;Species: Arabidopsis thaliana (mouse-ear cress)
C;Date: 02-Mar-2001 #sequence_revision 02-Mar-2001 #text_change 31-Mar-2001
C;Accession: H86206
R;Theologis, A.; Ecker, J.R.; Palm, C.J.; Federspiel, N.A.; Kaul, S.; White, O.; Al
Chin, C.W.; Chung, M.K.; Conn, L.; Conway, A.B.; Creasy, T.H.; Dewar
Ansen, N.F.; Hughes, B.; Huizar, L.
Nature 408, 816-820, 2000
A;Authors: Hunter, J.L.; Jenkins, J.J.; Johnson-Hopson, C.; Khan, S.; Khaykin, E.; Ki
C.A.; Li, J.H.; Li, Y.; Lin, X.; Liu, S.X.; Liu, Z.A.; Luos, J.S.; Maiti, R.; Marz
Rizzo, M.; Rooney, T.; Rowley, D.; Sarano, H.
A;Authors: Salzberg, S.L.; Schwarz, J.R.; Shinn, P.; Southwick, A.M.; Sun, H.; Tal
ker, M.; Wu, D.; Yu, G.; Fraser, C.M.; Venter, J.C.; Davis, R.W.
A;Title: Sequence and analysis of chromosome 1 of the Plant Arabidopsis.
A;Reference number: A86141; MUID:21016719
A;Accession: H86206
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-512 <STO>
A;Cross-references: GB:AE005172; NID:98954028; PIDN:AAF82202.1; GSPDB:GN00141
C;Genetics:
A;Map position: 1

Query Match 92.7%; Score 38; DB 2; Length 296;
Best Local Similarity 85.7%; Pred. No. 1.5;
Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
Db 275 KKRILHC 281

RESULT 4
S62624 biliverdin reductase (EC 1.3.1.24) - human
N;Alternate names: biliverdin IX-alpha reductase
C;Species: Homo sapiens (man)
C;Date: 28-Oct-1996 #sequence_revision 09-May-1997 #text_change 21-Jul-2000
C;Accession: S62644; S62622; S29736
R;Maines, M.D.; Polevsky, B.V.; Huaiq, T.J.; McCoubrey Jr., W.K.
Eur. J. Biochem. 235, 372-381, 1996
A;Title: Human biliverdin IX-alpha-reductase is a zinc-metalloprotein. Characterization
A;Accession number: S62622; MUID:96202961
A;Accession: S62624
A;Molecule type: mRNA
A;Residues: 1-296 <MAF>
A;Cross references: EMBL:X93086; PIDN:91246748; PIDN:CAA63635.1; PID:91246749
A;Accession: S62622
A;Molecule type: Protein
A;Residues: 3-24, X, 26-27, 'X', 29-35; 48-74; 228-234; 235-248 <MAF>
R;Maines, M.D.; Traksiel, G.M.
Arch. Biochem. Biophys. 300, 320-326, 1993
A;Title: Purification and characterization of human biliverdin reductase.
A;Accession: S29736
A;Molecule type: Protein
A;Accession number: S29736; MUID:93143333
A;Note: the sequence of peptide 1 from page 323 seems not to belong to this protein
C;Genetics:
C;Keywords: BVR
F;3-296/Product: biliverdin reductase IX-alpha #status experimental <MAF>

Query Match 92.7%; Score 38; DB 2; Length 296;
Best Local Similarity 85.7%; Pred. No. 1.5;
Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
Db 275 KKRILHC 281

RESULT 7
T01723 1-phosphatidylinositol-4-phosphate 5-kinase type II homolog - Arabidopsis thaliana
N;Alternate names: Protein A;IG002N01.9
C;Species: Arabidopsis thaliana (mouse-ear cress)
C;Date: 19-Feb-1999 #sequence_revision 19-Feb-1999 #text_change 24-Mar-1999
C;Accession: T01723
R;Scheet, P.; Maggi, L.

submitted to the EMBL Data Library, June 1997

A; Description: The sequence of *A. thaliana* IG002N01.
 A; Reference number: 214407
 A; Accession: T0173
 A; Status: translated from GB/EMBL/DDJB
 A; Molecule type: DNA
 A; Residues: 1-431 <SCH>
 A; Cross-references: EMBL:AF007269; NID:92191143
 A; Experimental source: cultivar Columbia
 C; Genetics:
 A; Map position: 4
 A; Introns: 40/2; 94/3; 161/3; 224/2; 255/1; 271/1; 303/1; 339/2
 A; Note: A_IG002N01.9

Query Match 80.5%; Score 33; DB 2; Length 189;
 Best Local Similarity 70.14%; Pred. No. 11; Indels 0; Gaps 0;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRMHC 7
 |||||
 Db 383 KKRIEHC 389

RESULT 8
 A38117 hypothetical protein 1 - *Anabaena* sp. insertion sequence IS895
 C;Species: *Anabaena* sp.
 C;Date: 24-Jul-1992 #sequence_revision 24-Jul-1992 #text_change 15-Oct-1999
 C;Accession: A38117
 R;Alam, J.; Vrba, J.M.; Cai, Y.; Martin, J.A.; Weislo, L.J.; Curtis, S.E.
 J; Bacteriol. 173: 5778-5783, 1991
 A;Title: Characterization of the IS895 family of insertion sequences from the cyanobacteria
 A;Reference number: A38117; MUID:91383700
 A;Accession: A38117
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 1-189 <ALA>
 A;Cross-references: GB:M67475; NID:9142026; PIDN:AAA98138.1; PID:9142027
 A;Experimental source: strain PCC 7120
 C;Genetics:
 A;Mobile element: insertion sequence IS895

Query Match 80.5%; Score 33; DB 2; Length 189;
 Best Local Similarity 70.14%; Pred. No. 11; Indels 0; Gaps 0;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRMHC 7
 |||||
 Db 171 KKRLXHC 177

RESULT 9
 T02486 hypothetical protein A-2929990 [imported] - *Arabidopsis thaliana*
 N;Alternate names: hypothetical protein F23F1.9
 C;Species: *Arabidopsis thaliana* (mouse-ear cress)
 C;Date: 05-Mar-1999 #sequence_revision 05-Mar-1999 #text_change 16-Feb-2001
 C;Accession: T02486; BB4703
 R;Rounseley, S.D.; Lin, X.; Ketchum, K.A.; Crosby, M.L.; Brandon, R.C.; Sykes, S.M.; Kaul, A;Description: *Arabidopsis thaliana* chromosome II BAC F23F1 genomic sequence.
 A;Reference number: Z14675
 A;Accession: T02486
 A;Status: translated from GB/EMBL/DDJB
 A;Molecule type: DNA
 A;Residues: 1-508 <ROU>
 A;Cross-references: EMBL:AC004680; NID:93420043; PID:93420052
 A;Experimental source: cultivar Columbia
 R;Lin, X.; Kaul, S.; Rounseley, S.D.; Shea, T.P.; Benito, M.I.; Town, C.D.; Fujii, C.Y.; M.; Koo, H.; Moffat, K.S.; Cronin, L.A.; Shen, M.; vanAken, S.B.; Umayam, L.; Talon, L.; euss, D.; Nierman, W.C.; White, O.; Eisen, J.A.; Salzberg, S.L.; Fraser, C.M.; Venter, C.

Nature 402, 761-768, 1999
 A;Title: Sequence and analysis of chromosome 2 of the plant *Arabidopsis thaliana*.
 A;Reference number: A84420; MUID:20083487
 A;Accession: B4703
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 1-308 <STO>
 A;Cross-references: GB:AB002093; NID:93420052; PIDN: AAC31853.1; GSPDB:GN00139
 C;Genetics:
 A;Gene: Atg29990; F23F1.9
 A;Map Position: 2
 A;Introns: 158/3; 230/1; 283/3; 305/3; 360/2; 398/3; 458/3
 C;Superfamily: NADH dehydrogenase

Query Match 80.5%; Score 33; DB 2; Length 508;
 Best Local Similarity 57.1%; Pred. No. 28; Mismatches 3; Indels 0; Gaps 0;
 Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRMHC 7
 |||||
 Db 236 KRRLEHC 242

RESULT 10
 S50490 hypothetical protein YERR32w - yeast (Saccharomyces cerevisiae)
 C;Species: *Saccharomyces cerevisiae*
 C;Date: 28-May-1993 #sequence_revision 24-Feb-1995 #text_change 23-Mar-2001
 C;Accession: S50490
 R;Dietrich, F.S.
 submitted to the EMBL Data Library December 1994
 A;Description: The sequence of *S. cerevisiae* cosmids 9537, 9581, 9495, 9867, and 1e
 A;Reference number: S50433
 A;Accession: S50490
 A;Molecule type: DNA
 A;Residues: 1-925 <DEI>
 A;Cross-references: EMBL:U18778; NID:9603592; PIDN: AAB64565.1; PID:9603624; MIPS:YI
 C;Genetics:
 A;Gene: SGD:FIR1
 A;Cross-references: SGD:S0000034; MIPS:YER032w
 A;Map position: 5R

Query Match 80.5%; Score 33; DB 2; Length 925;
 Best Local Similarity 71.4%; Pred. No. 48; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRMHC 7
 |||||
 Db 841 KKRLSHC 847

RESULT 11
 B64845 hypothetical protein b1028 - *Escherichia coli*
 C;Species: *Escherichia coli*
 C;Accession: B64845
 R;Blattner, F.R.; Plunkett III, G.; Bloch, C.A.; Perna, N.T.; Burland, V.; Riley, K.; Rose, D.J.; Mau, B.; Shao, Y.
 Science 277, 1453-1462, 1997
 A;Title: The complete genome sequence of *Escherichia coli* K-12.
 A;Reference number: A64720; MUID:97426617
 A;Accession: B64845
 A;Status: nucleic acid sequence not shown; translation not shown
 A;Molecule type: DNA
 A;Residues: 1-107 <BLAT>
 A;Cross-references: GB:U000936; NID:91787265; PIDN: AAC74113.1; PID:9176

Query Match 78.0%; Score 32; DB 2; Length 107;
 Best Local Similarity 57.1%; Pred. No. 11;
 Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

Qy 1 KKRIMHC 7
 Db 90 QQRYMHC 96

RESULT 12

D85651 hypothetical protein 21557 [imported] - Escherichia coli (strain O157:H7)
 C;Species: Escherichia coli
 C;Date: 16-Feb-2001 #sequence_revision 16-Feb-2001 #text_change 31-Mar-2001
 C;Accession: D85651
 R;Perna, N.T.; Plunkett III, G.; Burland, V.; Mau, B.; Glaser, J.D.; Rose, D.J.; Mayhew, L.; Grotbeck, E.J.; Davis, N.W.; Lim, A.; Dimalanta, E.; Potanousis, K.; Apodaca, Nature 409, 529-533, 2001
 A;Title: Genome sequence of enterohemorrhagic Escherichia coli O157:H7.
 A;Reference number: A85480; PMID:21074935; PMID:11206551
 A;Accession: D85651
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 1-107 <STO>
 A;Cross references: GB:AE005174; NID:912514431; PIDN:AAG55672.1; GSPDB:GN00145; UWGP:Z15
 A;Experimental source: strain O157:H7, substrate EDL933
 C;Genetics:
 A;Gene: Z1557
 C;Superfamily: Escherichia coli hypothetical protein b1028

Query Match 78.0%; Score 32; DB 2; Length 107;
 Best Local Similarity 57.1%; Pred. No. 11;
 Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

Qy 1 KKRIMHC 7
 Db 90 QQRYMHC 96

RESULT 13

I49668 binding protein - mouse
 C;Species: Mus musculus (house mouse)
 C;Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 16-Jul-1999
 C;Accession: I49668
 R;Hendrickson, B.A.; Zhang, W.; Craig, R.J.; Jin, Y.
 Gene 134, 271-275, 1993
 A;Title: Structural organization of the genes encoding human and murine FK506-binding protein
 A;Accession number: 149668; MUID:94085790
 A;Accession: I49668
 A;Status: preliminary; translated from GB/EMBL/DDBJ
 A;Molecule type: DNA
 A;Residues: 1-140 <RES>
 C;Cross references: GB:M77831; NID:9433782; PIDN:AAA37631.1; PID:9433783
 C;Genetics:
 A;Gene: Pkbp13
 A;Protein: 55.3; 93/2; 109/1; 121/1
 C;Superfamily: BKBP-type peptidylprolyl isomerase; BKBP-type peptidylprolyl isomerase hc
 F;47-94/Domain: BKBP-type peptidylprolyl isomerase homology <PPI>

Query Match 78.0%; Score 32; DB 2; Length 140;
 Best Local Similarity 71.4%; Pred. No. 14;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 KKRIMHC 7
 Db 34 KKRVDHc 40

RESULT 14

JC1365 FK506/rapamycin-binding protein FKBP13 precursor - human
 C;Species: Homo sapiens (man)
 C;Date: 30-Sep-1993 #sequence_revision 30-Sep-1993 #text_change 07-Aug-1998
 C;Accession: JC1365; A35602
 R;Dieleman, A.G.; Hawkin, A.; Craig, R.J.; Schreiber, S.L.; Griffin, C.A.
 Biochem. Biophys. Res. Commun. 189, 819-823, 1992
 A;Title: Chromosomal band assignments of the genes encoding human FKBP12 and FKBP13
 A;Reference number: JC1365; MUID:93112052
 A;Accession: JC1365
 A;Molecule type: DNA
 A;Residues: 1-142 <SDIL>
 R;Jin, Y.J.; Albert, M.W.; Lane, W.S.; Bierer, B.E.; Schreiber, S.L.; Burakoff, S.J.
 Proc. Natl. Acad. Sci. U.S.A. 88, 6677-6681, 1991
 A;Title: Molecular cloning of a membrane-associated human FK506- and rapamycin-bind
 A;Reference number: A39602; MUID:91319747
 A;Accession: A39602
 A;Molecule type: mRNA
 A;Residues: 1-20; 'S'; 23-142 <JIN>
 A;Cross-references: GB:M65128
 C;Genetics:
 A;Gene: GDB:FKBP2
 A;Cross-references: GDB:1133728; OMIM:186946
 A;Map position: 11q13.1-11q13.3
 A;Introns: 57; 95/2; 111/1; 123/1
 C;Superfamily: BKBP-type peptidylprolyl isomerase; BKBP-type peptidylprolyl
 C;Keywords: immunoregulation
 F;1-22/Domain: signal sequence #status predicted <SIG>
 F;23-142/Product: FK506/rapamycin-binding protein FKBP13 #status predicted <PP1>
 F;49-96/Domain: BKBP-type peptidylprolyl isomerase homology <PPI>

Query Match 78.0%; Score 32; DB 2; Length 142;
 Best Local Similarity 71.4%; Pred. No. 14;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 KKRIMHC 7
 Db 36 KKRVDHc 42

RESULT 15

B65086 hydrogenase (EC 1.18.99.1) 2 small chain - Escherichia coli
 C;Species: Escherichia coli
 C;Date: 12-Sep-1997 #sequence_revision 17-Sep-1997 #text_change 17-Mar-2000
 C;Accession: B65086; A55116
 R;Blattner, R.R.; Plunkett III, G.; Bloch, C.A.; Perna, N.T.; Burland, V.; Riley, M.
 A;Rose, D.J.; Mau, B.; Shao, Y.
 Science 277, 1453-1462, 1997
 A;Title: The complete genome sequence of Escherichia coli K-12.
 A;Reference number: A64720; MUID:9426657
 A;Accession: B65086
 A;Status: nucleic acid sequence not shown; translation not shown
 A;Molecule type: DNA
 A;Residues: 1-128 <BLAT>
 A;Cross-references: GB:AB000382; GB:U00096; NID:92367182; PIDN: AAC76032.1; PID:g178
 A;Experimental source: strain K-12, substrate MG1655
 R;Menon, N.K.; Chatelet, C.Y.; Devrathan, M.; Wendt, J.C.; Shanmugam, K.T.; Peck J; Bacteriol. 176, 4416-4423, 1994
 A;Title: Cloning, sequencing, and mutational analysis of the hyb operon encoding Es
 A;Reference number: A55516; MUID:94292472
 A;Accession: A55516
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 'V'; 2-328 <MEND>
 A;Cross-references: GB:U09177
 C;Genetics:
 A;Gene: hyA
 C;Superfamily: oxidoreductase
 C;Keywords: oxidoreductase
 F;105-163/Domain: ferredoxin 2[4Fe-4S] homology <FER-4>

Query Match 78.0%; Score 32; DB 2; Length 328;
Best Local Similarity 71.4%; Pred. No. 30;
Matches 5; Conservative 1; Mismatches 1; Indels 0;
Gaps 0;

QY 1 KKRTMHC 7
Db 109 KKQCMHC 115

Search completed: February 12, 2002, 11:51:40
Job time: 301 sec

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Perfect score:	41		
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Scoring table:	BLOSUM62		
GapPenalty:	10.0 , Gapext 0.5		
Searched:	522463 seqs, 74073290 residues		
Total number of hits satisfying chosen parameters:	522463		
Minimum DB seq length:	0		
Maximum DB seq length:	2000000000		
Post-processing:	Minimum Match 0% Maximum Match 100%		
	Listing first 45 summaries		
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			AC AAG28710;
			XX DT 17-OCT-2000 (first entry)
			XX DE Arabidopsis thaliana protein fragment SEQ ID NO: 34031.
			XX KW Protein identification; signal transduction pathway; metabolic pathway; hybridisation assay; generic mapping; gene expression control; promoter; terminatin sequence.
			XX KW Arabidopsis thaliana.
			XX OS EP1033405-A2.
			XX PN 06-SEP-2000.
			PD XX PF 25 - FEB - 2000; 2000EP-0301439.
			PF XX PR 25 - FEB - 1999; 99US-0121825.
			PR 14 - MAR - 1999; 99US-0123180.
			PR 15 - MAR - 1999; 99US-0125788.
			PR 16 - MAR - 1999; 99US-0126785.
			PR 17 - MAR - 1999; 99US-0127462.
			PR 18 - APR - 1999; 99US-018234.
			PR 19 - APR - 1999; 99US-0129845.
			PR 20 - APR - 1999; 99US-0130077.
			PR 21 - APR - 1999; 99US-0130449.

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query	Match	Length	DB ID	Description
1	36	87.8	285	21 AAG28710	Arabidopsis thalia Murine NGb1 isoform
2	34	82.9	94	21 AAY97243	Murine P371NG1 polypeptide
3	34	82.9	279	21 AAY97242	Arabidopsis thalia
4	33	80.5	508	21 AAG4146	Arabidopsis thalia
5	33	80.5	533	21 AAG4145	Arabidopsis thalia
6	33	80.5	564	21 AAG4144	Bovine FBP-13 immunoreactive protein
7	32	78.0	40	17 AAR35552	Human colon cancer
8	32	78.0	83	22 AAG76114	Human NGb1 isoform
9	32	78.0	94	21 AAY97245	Bovine FBP-13
10	32	78.0	99	13 AAR29797	Bovine secreted protein
11	32	78.0	104	21 AAGG3758	

PR 23-APR-1999; 99US-0130510.
 PR 28-APR-1999; 99US-0130891.
 PR 28-APR-1999; 99US-0131449.
 PR 30-APR-1999; 99US-0132048.
 PR 30-APR-1999; 99US-0132407.
 PR 04-MAY-1999; 99US-0132484.
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 PR 04-JUN-1999; 99US-0137502.
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 PR 14-OCT-1999; 99US-0159340.
 PR 14-OCT-1999; 99US-0159331.

Query	Match	Score	DB	Length	21;	285;	
			Pred	No.	Mismatches	0;	Gaps
R	R	99US-0159637.					
R	R	99US-0159638.					
R	R	99US-0159584.					
R	R	99US-0160741.					
R	R	99US-0160767.					
R	R	99US-0160768.					
R	R	99US-0160770.					
R	R	99US-0160814.					
R	R	99US-0160815.					
R	R	99US-0160980.					
R	R	99US-0160981.					
R	R	99US-0160989.					
R	R	99US-0161404.					
R	R	99US-0161405.					
R	R	99US-0161406.					
R	R	99US-0161359.					
R	R	99US-0161360.					
R	R	99US-0161361.					
R	R	99US-0161920.					
R	R	99US-0161992.					
R	R	99US-0161993.					
R	R	99US-0162142.					
Y	Y	1. KKRIMHC 7 : 13 krikli 19					

ING1. Functional cooperation between ING1 and p53 suggested that ING1 encoded a tumour suppressor protein that functioned within the p53 pathway. This data suggested a possible role for ING1 in head and neck cancers and chromosomal location of the ING1 placed it within a region that is frequently rearranged in head and neck cancers. Large scale analysis of tumours involving ING1 has not revealed mutations in ING1 nor significant variations in its expression suggesting that ING1 was not a useful gene to study in cancer etiology. However, alternative initiation exons of the ing1 gene, each having their own promoter have been discovered.

Expression of one promoter (la) produces a protein having an identical C-terminal fragment to ING1 but an additional 104 N-terminal amino acids. The newly discovered protein has been designated p31ING1 (Wild type; p33ING1). P37ING1 has the characteristics of an oncogene. When overexpressed in cells (even those expressing wild type p53) P37ING1 is able to cause proliferation or transformation of those cells. Thus detecting a nucleic acid encoding exon 1b of ing1 by hybridisation with an isolated nucleic acid having the sequence of exon 1b of ing1 or its antisense sequence can identify individuals expressing the oncogenic form of ing1. Novel peptide sequences taken from the 104 N-terminal peptide of p37ING1 can also be used to raise antibodies that can also be used in detection methods for the P37ING1 variant. The polypeptides may be useful in gene therapy for treatment of cell proliferation disorders, especially cancers and for diagnosing and studying cancers.

Sequence	94 AA;
Query Match	82 %;
Best Local Similarity	57.1%;
Matches	Pred. No. 12;
4 ; Conservative	Mismatches 3;
QY	Indels 0;
1 KKRIMHC 7	Gaps 0
Db	
67 krrvrc 73	

RESULT

Query Match	Score 34;	DB 21;	Length 94;
XX			
AC			
XX			
DT			
19-DECC-2000	(first entry)		
XX			
DE			
Murine P37ING1 polypeptide.			
XX			
KW			
ING1; tumour; p33ING1; p37ING1; oncogene; gene therapy; diagnosis;			
KW			
KW			
proliferation disorder; transformation; transformed cell; mouse.			
XX			
OS			
Mus musculus.			
XX			
PN			
WO200046370-A1.			
XX			
PR			
04 - FEB - 1999;			
XX			
(UNII) UNIV ILLINOIS FOUND.			
PA			
PI			
Gudikov A, Zeremski M, Gurova KV, Grigorian IA;			
XX			
DR			
WPI ; 2000-491278/43 .			
DR			
N-PSDB; AAA53730.			
XX			
PT			
Detecting nucleic acid encoding exon 1b of ing1, useful for diagnosing and treating cancer, comprises contacting sample with isolated nucleic			

acid comprising sequence of exon 1b and detecting hybridized products disclosure. Fig. 12. 13400. English

Mutations in or loss of the p53 gene occur in more than 50% of human tumours and tumour cell lines, but functional inactivation of the p53 pathway occurs in much larger proportion of tumours. In many cases the mechanism of functional inactivation of the p53 gene remains unknown but p53 has been found to act in cooperation with ING1. Functional cooperation between ING1 and p53 suggested that ING1 encoded a tumour suppressor protein that functioned within the p53 pathway. This data suggested a possible role for ING1 in head and neck cancers and chromosomal location of the ING1 placed it within a region that is frequently rearranged in head and neck cancers. Large scale analysis of tumours involving ING1 has not revealed mutations in ING1 nor significant variations in its expression suggesting that ING1 was not a useful gene to study in cancer etiology. However, alternative initiation exons of the ing1 gene, each having their own promoter have been discovered. Expression of one promoter (1a) produces a protein identical to ING1. Expression of a second promoter (1b) produces a protein having an identical C-terminal fragment to ING1 but an additional 104 N-terminal amino acids. The newly discovered protein has been designated P37ING1 (wild type; p37ING1). P37ING1 has the characteristics of an oncogene. When overexpressed in cells (even those expressing wild type p53) P37ING1 is able to cause proliferation or transformation of those cells. Thus detecting a nucleic acid encoding exon 1b of ing1 by hybridisation with an isolated nucleic acid having the sequence of exon 1b of ing1 or its antisense sequence can identify individuals expressing the oncogenic form of ing1. Novel peptide sequences taken from the 104 N-terminal Peptide of P37ING1 can also be used to raise antibodies that can also be used in detection methods for the p37ING1 variant. The polypeptides may be used in gene therapy for treatment of cell proliferation disorders, especially cancers and for diagnosing and screening cancers.

Sequence 279 AA; Query Match 82.9%; Best Local Similarity 57.1%; Score 34; DB 21; Pred. No. 34; Length 279;

Query	Match	Best	Local	Similarity	4;	Conser-
		Matches				ves
Y	1	KKRIMHC	7			
		1 : : :				

RESULT 4

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06-APR-1999 PPR 99US-0128334.
08-APR-1999 PPR 99US-0128114.
16-APR-1999 PPR 99US-0129455.
19-APR-1999 PPR 99US-0130000.
21-APR-1999 PPR 99US-0130459.
23-APR-1999 PPR 99US-0130510.
23-APR-1999 PPR 99US-0130891.
28-APR-1999 PPR 99US-0131149.
30-APR-1999 PPR 99US-0132248.
01-MAY-1999 PPR 99US-0132463.
04-MAY-1999 PPR 99US-0132484.
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Query Match 80.5%; Score 33; DB 21; Length 508;
 Best Local Similarity 57.1%; Pred. No. 95; Mismatches 0; Indels 0; Gaps 0;

Qy 1 KKRIMHC 7
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 Db 236 krrimhc 242

RESULT 5
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 ID XX
 AC XX
 DT XX
 DT 18-OCT-2000 (first entry)
 DE XX
 DE Arabidopsis thaliana protein fragment SEQ ID NO: 60769.
 KW XX
 KW Protein identification; signal transduction pathway; metabolic pathway;
 KW hybridisation assay; genetic mapping; gene expression control; promoter;
 KW termination sequence.
 XX XX
 OS OS
 XX XX
 PN EP1033405-A2.
 XX XX
 PD 06-SEP-2000.
 XX XX
 PP 25-FEB-2000; 2000EP-0301439.
 XX XX
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 PR 05-MAR-1999; 99US-0123180.
 PR 09-MAR-1999; 99US-0123548.
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 PR 28-OCT-1999; 99US-0161992.
 PR 28-OCT-1999; 99US-0161993.
 PR 29-OCT-1999; 99US-0162142.

Query Match 80.5%; Score 33; DB 21; Length 533;
 Best Local Similarity 57.1%; Pred. NO. 99; Gaps 0;
 Matches 4; Conservative 3; Mismatches 0; Indels 0;

Qy 1 KKRIMHC 7
 Db 261 krrlhc 267

RESULT 6
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 ID AAG48144
 AC AAG48144;
 XX DT 18-OCT-2000 (first entry)
 DE Arabidopsis thaliana protein fragment SEQ ID NO: 60768.
 XX KW Protein identification; signal transduction pathway; metabolic pathway;
 KW hybridisation assay; genetic mapping; gene expression control; promoter;
 KW termination sequence.
 XX OS Arabidopsis thaliana.
 XX PN EP103405-A2.
 XX PD 06-SEP-2000.
 XX PR 25-FEB-2000; 2000EP-0301439.

PR 30-APR-1999; 99US-0132407.
 PR 04-MAY-1999; 99US-0132484.
 PR 05-MAY-1999; 99US-0132485.
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 PR 11-MAY-1999; 99US-0134256.
 PR 14-MAY-1999; 99US-0134218.
 PR 14-MAY-1999; 99US-0134219.
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 PR 14-MAY-1999; 99US-0134370.
 PR 18-MAY-1999; 99US-0134768.
 PR 19-MAY-1999; 99US-0134911.
 PR 20-MAY-1999; 99US-0134912.
 PR 21-MAY-1999; 99US-0135533.
 PR 24-MAY-1999; 99US-0135629.
 PR 25-MAY-1999; 99US-0136021.
 PR 27-MAY-1999; 99US-0136592.
 PR 28-MAY-1999; 99US-0136782.
 PR 01-JUN-1999; 99US-0137222.
 PR 03-JUN-1999; 99US-0137558.
 PR 04-JUN-1999; 99US-0137512.
 PR 07-JUN-1999; 99US-0137724.
 PR 08-JUN-1999; 99US-0138094.
 PR 10-JUN-1999; 99US-0138540.
 PR 10-JUN-1999; 99US-0138847.
 PR 14-JUN-1999; 99US-0139119.
 PR 16-JUN-1999; 99US-0139452.
 PR 16-JUN-1999; 99US-0139453.
 PR 17-JUN-1999; 99US-0139452.
 PR 18-JUN-1999; 99US-0139444.
 PR 18-JUN-1999; 99US-0139455.
 PR 18-JUN-1999; 99US-0139456.
 PR 18-JUN-1999; 99US-0139457.
 PR 18-JUN-1999; 99US-0139458.
 PR 18-JUN-1999; 99US-0139459.
 PR 18-JUN-1999; 99US-0139460.
 PR 18-JUN-1999; 99US-0139461.
 PR 18-JUN-1999; 99US-0139462.
 PR 18-JUN-1999; 99US-0139463.
 PR 18-JUN-1999; 99US-0139750.
 PR 18-JUN-1999; 99US-0139763.
 PR 21-JUN-1999; 99US-0139817.
 PR 22-JUN-1999; 99US-0139899.
 PR 23-JUN-1999; 99US-0140352.
 PR 23-JUN-1999; 99US-0140354.
 PR 24-JUN-1999; 99US-0140695.
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 PR 30-JUN-1999; 99US-0141281.
 PR 01-JUL-1999; 99US-0141842.
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 PR 02-JUL-1999; 99US-0142055.
 PR 06-JUL-1999; 99US-0142390.
 PR 08-JUL-1999; 99US-012803.
 PR 09-JUL-1999; 99US-0142977.
 PR 12-JUL-1999; 99US-0143542.
 PR 13-JUL-1999; 99US-0143624.
 PR 14-JUL-1999; 99US-0144005.
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PR 21-JUL-1999; 99US-0145088; PR 21-OCT-1999; 99US-0160767.
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 PR 27-JUL-1999; 99US-0145919; PR 26-OCT-1999; 99US-0161359.
 PR 28-JUL-1999; 99US-0145951; PR 26-OCT-1999; 99US-0161360.
 PR 02-AUG-1999; 99US-0146386; PR 26-OCT-1999; 99US-0161361.
 PR 02-AUG-1999; 99US-0146388; PR 28-OCT-1999; 99US-016120.
 PR 02-AUG-1999; 99US-0146389; PR 28-OCT-1999; 99US-0161992.
 PR 03-AUG-1999; 99US-0147204; PR 28-OCT-1999; 99US-0161993.
 PR 04-AUG-1999; 99US-0147302; PR 29-OCT-1999; 99US-0161442.
 PR 05-AUG-1999; 99US-0147260; PR 29-OCT-1999; 99US-0161442.

Query Match Score 33; DB 21; Length 564;
 Best Local Similarity 57.1%; Pred: NO. 1e+02; Mismatches 0; Indels 0; Gaps 0;

Qy 1 KKRMHC 7
 Db 292 krrlhc 298

RESULT 7
 AAR93552
 ID AAR93552 standard; Peptide; 40 AA.
 XX
 AC AAR93552;
 XX
 25-JUN-1996 (first entry)
 DT DT
 XX
 DE Bovine FKBP-13 immunophilin N-terminal peptide.
 XX
 KW FKBP-13; immunophilin; FK506; rapamycin; rheumatoid arthritis;
 KW diabetes; organ transplant; graft versus host disease;
 KW immunosuppressant.
 XX
 Bos taurus.
 OS OS
 PN US5498597-A.
 XX
 (DAND) DANA FARBER CANCER INST INC.
 XX
 PD 12-MAR-1996.
 XX
 PR 20-AUG-1999; 99US-0149723; PR 20-AUG-1999; 99US-0149565.
 PR 20-AUG-1999; 99US-0148684; PR 16-AUG-1999; 99US-0149368.
 PR 17-AUG-1999; 99US-0149175; PR 18-AUG-1999; 99US-0149426.
 PR 20-AUG-1999; 99US-0149722; PR 20-AUG-1999; 99US-0149723.
 PR 20-AUG-1999; 99US-0149929; PR 23-AUG-1999; 99US-0149902.
 PR 23-AUG-1999; 99US-0149930; PR 25-AUG-1999; 99US-0150566.
 PR 26-AUG-1999; 99US-0150884; PR 27-AUG-1999; 99US-0151065.
 PR 27-AUG-1999; 99US-0151066; PR 27-AUG-1999; 99US-0151080.
 PR 30-AUG-1999; 99US-0151301; PR 31-AUG-1999; 99US-0151436.
 PR 01-SEP-1999; 99US-0151930; PR 01-SEP-1999; 99US-0151930.
 PR 07-SEP-1999; 99US-0152363; PR 10-SEP-1999; 99US-0153070.
 PR 13-SEP-1999; 99US-0153758; PR 15-SEP-1999; 99US-0154018.
 PR 16-SEP-1999; 99US-0154039; PR 20-SEP-1999; 99US-0154779.
 PR 22-SEP-1999; 99US-0155139; PR 22-SEP-1999; 99US-0155263.
 PR 24-SEP-1999; 99US-0155659; PR 28-SEP-1999; 99US-0158232.
 PR 28-SEP-1999; 99US-0156596; PR 29-SEP-1999; 99US-0158369.
 PR 12-OCT-1999; 99US-0156596; PR 12-OCT-1999; 99US-0158369.
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 PR 13-OCT-1999; 99US-0159295; PR 13-OCT-1999; 99US-0159295.
 PR 14-OCT-1999; 99US-0159329; PR 14-OCT-1999; 99US-0159329.
 PR 14-OCT-1999; 99US-0159330; PR 14-OCT-1999; 99US-0159331.
 PR 14-OCT-1999; 99US-0159637; PR 14-OCT-1999; 99US-0159638.
 PR 18-OCT-1999; 99US-0159584; PR 18-OCT-1999; 99US-0160741.

CC This sequence encoding the bovine FKBP-13 N-terminal sequence
 CC corresponds to the N-terminal sequence of human FKBP-13. FKBP-13
 CC may be used for identifying immunosuppressant drugs, and may be
 CC used in combination with immunosuppressant drugs for therapeutic
 CC purposes in the treatment of autoimmune diseases e.g. rheumatoid
 CC arthritis and type-I diabetes, organ transplant and graft versus
 CC host disease. The recombinant form of the protein could be
 CC potentially smaller and therefore easier to introduce into cells
 CC than intact FKBP-13.

CC The polypeptides may be useful in gene therapy for treatment of cell
 CC proliferation disorders, especially cancers and for diagnosing and
 CC studying cancers.

XX RESULT 11
 SQ Sequence 94 AA;
 Query Match 78.0%; Score 32; DB 21; Length 94;
 Best Local Similarity 57.1%; Pred. No. 30;
 Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 KKRIMHC 7
 Db 67 krrimhc 73

XX AAG03758 standard; Protein: 104 AA.
 ID AAG03758
 AC AAG03758;
 XX DT 06-OCT-2000 (first entry)
 DE Human secreted protein, SEQ ID NO: 7839.
 XX Human; 5' EST; expressed sequence tag; secreted protein; cDNA isolation;
 KW gene therapy; chromosome mapping.
 XX OS Homo sapiens.
 XX PN EP1031401-A2.
 AC PD 06-SEP-2000.
 XX PF 21-FEB-2000; 2000EP-0200610.
 XX PR 26-FEB-1999; 99US-0122487.
 XX PA (GEST) GENSET.
 XX PI Dumas Milne Edwards J, Ducleit A, Giordano J;
 XX WPI; 2000-500381/45.
 DR N-PSDB, AAC03764.
 XX PT New nucleic acid that is a 5' expressed sequence tag (5' EST) for
 PT obtaining cDNAs and genomic DNAs that correspond to 5'ESTs and for
 PT diagnostic, forensic, gene therapy and chromosome mapping procedures -
 PS Claim 13; SEQ ID 7839: 71pp + CD-ROM; English.
 XX PT The present sequence is a polypeptide encoded by one of a large number
 PT of 5' ESTs derived from mRNAs encoding secreted proteins. The 5' ESTs
 CC were prepared from total human RNAs or polyA+ RNAs derived from 30
 CC different tissues. EST sequences usually correspond mainly to the 3'
 CC untranslated region (UTR) of the mRNA because they are often obtained
 CC from oligo-dT primed cDNA libraries. Such ESTs are not well suited for
 CC isolating cDNA sequences derived from the 5' ends of mRNAs and even in
 CC those cases where longer cDNA sequences have been obtained, the full 5'
 CC UTR is rarely included. 5' ESTs are derived from mRNAs with intact 5'
 CC ends and can therefore be used to obtain full length cDNAs and genomic
 CC DNAs. 5' ESTs are also used in diagnostic, forensic, gene therapy and
 CC chromosome mapping procedures. They are used to obtain upstream
 CC regulatory sequences and to design expression and secretion vectors.
 XX SQ Sequence 104 AA;

XX Query Match 78.0%; Score 32; DB 21; Length 104;
 Best Local Similarity 71.4%; Pred. No. 33;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 PS Disclosure: Fig 1; 30pp; English.
 XX Qy 1 KKRIMHC 7
 Db 36 kkrvhc 42

XX RESULT 12
 SQ Sequence 99 AA;
 Query Match 78.0%; Score 32; DB 13; Length 99;
 Best Local Similarity 71.4%; Pred. No. 32;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 PS Qy 1 KKRIMHC 7
 Db 14 kkrvhc 20

XX Human; p33-ING1 protein; growth regulation; apoptosis; DNA damage;
 KW inhibition; anchorage independent growth; cytotoxic drug; cancer;
 KW transcriptional activation; immortal cell line; p28-ING1 protein.
 XX OS Homo sapiens.
 XX FH Key
 FT Misc-Difference 59-269 Location/Qualifiers
 /note= "p26-ING1 fragment"
 XX PN WO9916790-A1.
 PD 08-APR-1999.
 XX PF 24-SEP-1998; 98WO-US18179.
 XX PR 14-JAN-1998; 98US-0006783.
 PR 26-SEP-1997; 97US-0060138.
 XX PA (UNIV) UNIV ILLINOIS BOARD OF TRUSTEES.
 PA (UNIV) UNIV TECHNOLOGIES INT INC.
 PI Garkavtsev I, Gudkov A, Riabowol K;
 DR WPI: 1999-2633685/22.
 XX PT Use of p33-ING1 Peptides
 XX PS Example 8; Page 61; 64pp; English.
 XX This is the amino acid sequence of the human p28-ING1 protein.
 CC used in the method of the invention, involving the human p33-ING1
 protein. The ING1 gene encodes p33-ING1 which can be used to
 CC modulate the activity of, isolate or detect p53. Expression of the
 CC ING1 and p53 genes in a mammalian cell results in normal growth
 CC regulation and anchorage-dependent growth and apoptosis as a response
 CC expression of either gene results in a loss of cellular growth
 CC control, anchorage independent growth, inhibition of apoptosis
 CC and resistance to radiation and cytotoxic drugs. The p33-ING1 is a
 CC component of the p53 signalling pathway that cooperates with p53 in
 CC negative regulation of cell proliferation by modulating p53 dependent
 CC transcriptional activation. Biological function of p53 signalling
 CC pathway can therefore be regulated (both enhanced or suppressed) by
 CC modulating p33-ING1 activity. The modulation of p33-ING1 activity can
 CC be used for the stimulation or restoration of the p53 pathway to
 CC defend sensitive tissues from genotoxic stress or for the generation
 CC of immortal cell lines.
 XX SQ Sequence 128 AA;

Query Match Score 32; DB 20; Length 128;
 Best Local Similarity 57.1%; Pred. No. 41;
 Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 Db 67 krrmihc 73

RESULT 13
 AAB87656 ID AAB87656 standard; protein; 137 AA.
 XX AC AAB87656;
 XX DT 15-MAY-2001 (first entry)
 XX DE Bovine mammary tissue derived protein #47.
 XX

XX KW Bovine; mammary gland; cancer; tumour; angiogenesis.
 XX OS Bos taurus.
 XX PN WO200114553-A1.
 XX PD 01-MAR-2001.
 XX PF 23-AUG-2000; 2000WO-NZ00166.
 XX PR 23-AUG-1999; 99US-0150330.
 XX PA (GENE-) GENESIS RES & DEV CORP LTD.
 PA (NZPA-) NEW ZEALAND PASTORAL AGRIC RES INST LTD.
 XX PI Havukkala IJ, Gleen M, Grigor MR, Molenaar AJ;
 XX DR WPI; 2001-226619/23.
 XX PT New polypeptides and polynucleotides encoding the polypeptides, which
 PT are expressed in bovine mammary gland tissue, useful for stimulating
 PT mammary gland growth or function, or inducing differentiation of milk
 PT producing cells.
 XX C1aim 11; Page_80; 97PP; English.
 XX PS Sequence 137 AA;
 CC The present invention relates to proteins derived from bovine
 CC mammary cells. The invention is useful for stimulating the
 CC bovine mammary gland cell growth and function, inhibiting the
 CC growth of various mammary gland cancer cells, inhibiting
 CC angiogenesis and vascularization of tumours, or modulating
 CC the growth of blood vessels in a mammal.
 XX SQ Sequence 137 AA;

Query Match Score 32; DB 22; Length 137;
 Best Local Similarity 71.4%; Pred. No. 43;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 |||:
 Db 43 kkrvihc 49

RESULT 14
 AAR93551 ID AAR93551 standard; protein; 141 AA.
 XX AC AAR93551;
 XX DT 25-JUN-1996 (first entry)
 XX DE Human FKBP-13 immunophilin protein.
 XX FKBP-13; immunophilin; FK506; rapamycin; rheumatoid arthritis;
 KW matches; organ transplant; graft versus host disease;
 KW immunosuppressant.
 XX OS Homo sapiens.

Key Peptide

Location/Qualifiers
 1.:21
 /label= sig-peptide

Misc-difference 22..61
 FT /note= "corresponds to bovine thymus FKBP-13 N-
 terminal sequence"

Misc-difference 117..120
 FT /note= "endoplasmic reticulum retention sequence"

RESULT 13
 AAB87656 ID AAB87656 standard; protein; 137 AA.
 XX AC AAB87656;
 XX DT 15-MAY-2001 (first entry)
 XX DE Bovine mammary tissue derived protein #47.
 XX

US5498597-A.
 XX PN
 PD 12-MAR-1996.

XX 17-JAN-1992; 92US-0822966.
 XX PR 17-JAN-1992; 92US-0822966.
 XX PA (DAND) DANA FARBER CANCER INST INC.
 XX PA (HARD) HARVARD COLLEGE.
 XX PI Bierer BE, Burakoff SJ, Schreiber SL;
 XX WPI: 1996-159713/16.
 XX DR N-PSDB: AAT18037.
 PT Purified mammalian FKBP-13 polypeptide capable of binding FK506
 PT useful for identifying and studying immunosuppressant drugs
 XX PS Claim 1; Fig 1; 12PP; English.
 XX The FKBP-13 protein may be used for identifying immunosuppressant
 CC drugs, and may be used in combination with immunosuppressant drugs
 CC for therapeutic purposes in the treatment of autoimmune diseases e.g.
 CC rheumatoid arthritis and type-I diabetes, organ transplant and
 CC graft versus host disease. The recombinant form of the protein
 CC could be potentially smaller and therefore easier to introduce
 CC into cells than intact FKBP-13.
 XX Sequence 141 AA;

Query Match Score 32; DB 21; Length 141;
 Best Local Similarity 78.0%; Pred. No. 45;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 KKRIMHC 7
 |||:|||
 Db 35 kkrvdhc 41

RESULT 15
 AAB53247
 ID AAB53247 Standard; Protein: 141 AA.
 XX
 AC AAB53247;
 XX DT 09-MAR-2001 (first entry)
 XX DE Human colon cancer antigen protein sequence SEQ ID NO:787.
 XX KW Human; colon cancer; colon cancer antigen; diagnosis; detection;
 KW identification; cytosolic; cardiaoactive; neuroprotective; vulnerability;
 KW immunomodulatory; muscular; gynecological; gastrointestinal;
 KW nephrotropic; antiinfective; antibacterial; gene therapy; wound;
 KW neural disorder; immune system disorder; muscular disorder;
 KW reproductive disorder; gastrointestinal disorder; renal disorder;
 KW infectious disease; cardiovascular disorder.
 XX OS Homo sapiens.
 XX PN WO200055551-A1.
 XX PD 21-SEP-2000.
 XX PF 08-MAR-2000; 2000WO-US5883.
 XX PR 12-MAR-1999; 99US-0124270.
 XX PA (HUMA-) HUMAN GENOME SCI INC.
 XX PI Rosen CA, Ruben SM;
 XX DR WPI: 2000-587534/55.
 XX N-PSDB: AAC98004.

GenCore version 4.5
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OM protein - protein search, using sw model

Run on:

February 12, 2002, 12:03:25 ; Search time 94.82 Seconds
(without alignments)

12.341 Million cell updates/sec

Title: US-09-606-129A-19

Perfect score: 46

Sequence: 1 QKLCHOKK 8

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 473505 seqs, 146272329 residues

Total number of hits satisfying chosen parameters: 473505

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SPTREMBL_17:*

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1: sp_archaea;*
2: sp_bacteria;*
3: sp_fungi;*
sp_human;*
4: sp_invertebrate;*
5: sp_mammal;*
6: sp_micr;*
7: sp_organelle;*
8: sp_phage;*
9: sp_plant;*
10: sp_rapid;*
11: sp_rapid;*
12: sp_virus;*
13: sp_vertebrate;*
14: sp_unclassified;*

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query	Match Length	DB ID	Description
1	37	80.4	636	12	O90142
2	37	80.4	1063	12	Q9J844
3	35	76.1	295	11	Q9CY64
4	35	76.1	303	11	Q9DD21
5	35	76.1	420	10	O22019
6	34	73.9	208	5	Q94164
7	34	73.9	322	4	Q9BSM2
8	34	73.9	409	11	Q9D3R6
9	34	73.9	522	4	Q9H8F4
10	34	73.9	547	3	O74308
11	34	73.9	790	4	Q9NV9
12	34	73.9	1188	5	O9E143
13	34	73.9	1295	3	O13348
14	34	73.9	1473	11	Q63625
15	34	73.9	1654	4	Q9PY6
16	33	73.9	261	4	Q9H1N6
17	33	71.7	263	1	Q9YDU2
18	33	71.7	266	1	P96301
19	33	71.7	273	12	Q9PYW7

20	33	71.7	445	10	Q9BPL5
21	33	71.7	566	8	Q9mt29
22	33	71.7	566	4	Q9HZSO
23	33	71.7	588	4	Q9UJU1
24	33	71.7	673	11	Q9QV74
25	33	71.7	773	10	Q80631
26	33	71.7	1247	5	Q17461
27	32	69.6	87	5	Q95N6
28	32	69.6	163	2	Q9EG2
29	32	69.6	312	11	Q9CSE2
30	32	69.6	336	4	Q9495
31	32	69.6	380	5	Q21866
32	32	69.6	397	4	Q9h497
33	32	69.6	397	4	Q9h5E7
34	32	69.6	422	10	Q9SSP1
35	32	69.6	462	5	Q9UDB3
36	32	69.6	465	5	Q9X94
37	32	69.6	474	10	Q8588
38	32	69.6	548	20	Q20367
39	32	69.6	590	2	Q45490
40	32	69.6	628	12	Q90139
41	32	69.6	828	11	Q9q207
42	32	69.6	828	11	Q9d616
43	32	69.6	843	12	Q934E8
44	32	69.6	864	10	Q9C987
45	32	69.6	920	11	Q9JUK7

ALIGNMENTS

RESTLT	1	PRELIMINARY;	PRT;	636 AA.
ID	O90142	ID	O90142	PRELIMINARY;
AC	O90142	AC	O90142	PRELIMINARY;
DT	01-NOV-1998	DT	01-NOV-1998	(TREMBLE1. 08, Created)
DT	01-JUN-2001	DT	01-JUN-2001	(TREMBLE1. 08, Last sequence update)
DT	01-JUN-2001	DT	01-JUN-2001	(TREMBLE1. 17, Last annotation update)
DE	DNA POLYMERASE (FRAGMENT).	DE	DNA POLYMERASE (FRAGMENT).	
DN	DNA POLY.	DN	DNA POLY.	
OS	spodoptera exigua nucleopolyhedrovirus	OS	spodoptera exigua nucleopolyhedrovirus	
OC	dsDNA viruses, no RNA stage; Baculoviridae;	OC	Nucleopolyhedrovirus,	
OC	Nucleopolyhedrovirus.	OC	Nucleopolyhedrovirus.	
NCBI_TaxID	10454;	NCBI_TaxID	10454;	
RN	[1]	RN	SEQUENCE FROM N.A.	
RP	RP61868; AAC3749.1;	RP	SEQUENCE FROM N.A.	
RA	Burach D.M., Kumar C.A., Zaias A., Liang B., Tribe D.E.;	RA	Group II Nucleopolyhedrovirus Subgroups Revealed by Phylogenetic Analysis of Polymerase Gene Sequences.;	
RT	"	RT	"	
RT	Submitted (MAY-1998) to the EMBL/GenBank/DDBJ databases.	RT	Submitted (MAY-1998) to the EMBL/GenBank/DDBJ databases.	
RL	-1 - CATALYTIC ACTIVITY: N DEOXYNUCLEOSIDE TRIPHOSPHATE = N	RL	-1 - CATALYTIC ACTIVITY: N DEOXYNUCLEOSIDE TRIPHOSPHATE = N	
CC	PYROPHOSPHATE + DNA(N).	CC	PYROPHOSPHATE + DNA(N).	
CC	-1- SIMILARITY: BELONGS TO DNA POLYMERASE TYPE-B FAMILY.	CC	-1- SIMILARITY: BELONGS TO DNA POLYMERASE TYPE-B FAMILY.	
DR	EMBL; AF061868; AAC3749.1;	DR	EMBL; AF061868; AAC3749.1;	
DR	InterPro; IPR020504; DNA-pol_B.	DR	InterPro; IPR020504; DNA-pol_B.	
Pfam	PF0136; DNA-pol_B_2.	Pfam	PF0136; DNA-pol_B_2.	
PRINTS	PRINTS; PRO0106; DNAPOLB.	PRINTS	PRINTS; PRO0106; DNAPOLB.	
SMART	SMART; SMD0485; DNApolB.	SMART	SMART; SMD0485; DNApolB.	
DR	DNA replication; DNA-binding; DNA-directed DNA polymerase.	DR	DNA replication; DNA-binding; DNA-directed DNA polymerase.	
KW		KW		
FT	NON_TER 1	FT	NON_TER 1	
FT	NON_TER 1	FT	NON_TER 1	
SQ	SEQUENCE 636 AA; 73836 MW;	SQ	SEQUENCE 636 AA; 73836 MW;	
QY	Query Match 80.4%; Score 37; DB 12; Length 636;	QY	Query Match 80.4%; Score 37; DB 12; Length 636;	
	Best Local Similarity 85.7%; Pred. No. 12; Mismatches 6; Conservative 1; Indels 0; Gaps 0;		Best Local Similarity 85.7%; Pred. No. 12; Mismatches 6; Conservative 1; Indels 0; Gaps 0;	
Db	392 KLCHQK8	Db	392 KLCHQK8	

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AC	022019							
DT	01-JAN-1998 (TREMBLrel. 05, Created)							
DT	01-JAN-1998 (TREMBLrel. 05, Last sequence update)							
DT	01-JUN-2001 (TREMBLrel. 17, Last annotation update)							
DE	ORF420							
OS	Cyanidioschyzon merolae.							
OC	Eukaryota; Rhodophyta; Bangiophyceae; Cyanidioschyzon.							
OX	NCBI_TAXID=45137;							
RN	[1]	SEQUENCE FROM N.A.						
RP								
RA	Ohta N;							
RL	J. Plant Res. 110:235-245(1997).							
DR	D63675; BAA22815.1;							
DR	Mendel: 23095; Cyame-3036; 23095.							
DR	InterPro: IPR000178; IP2;							
DR	InterPro: IPR001950; SU11.							
DR	Pfam: PF02131; IF2;							
DR	ProDom: PD186100; IF2;							
DR	PS00118; SU11;							
DR	PROSITE: PS00118; SU11; UNKNOWN_1;							
SQ	SEQUENCE 420 AA; 47691 MW; A6CAE107B24B4E19 CRC64;							
RESULT	6							
Q94164	Q94164	PRELIMINARY;	PRT;	208 AA.				
AC	Q94164;							
DT	01-FEB-1997 (TREMBLrel. 02, Created)							
DT	01-FEB-1997 (TREMBLrel. 02, Last sequence update)							
DT	01-JUN-2001 (TREMBLrel. 17, Last annotation update)							
DE	SIMILARITY TO BPTI/KUNITZ INHIBITOR DOMAIN.							
GN	C10GB 2.							
OS	Caenorhabditis elegans.							
OC	Rhabditidae; Nematoda; Chromadorea; Rhabditida; Rhabditoidae;							
OC	Rhabditidae; Peledorinae; Caenorhabditida.							
OX	NCBI_TAXID=6239;							
RN	[1]	SEQUENCE FROM N.A.						
RC	SEQUENCE=BRISTOL N2;							
RX	Medline=94150718; PubMed=7906398;							
RA	Wilson R., Ainscough R., Anderson K., Baynes C., Berks M., Coulson A., Bonfield J., Burton J., Connell M., Copsey T., Cooper J., Coulson A., Coxton M., Dear S., Du Z., Durbin R., Pavelllo A., Fulton L., Gardner A., Green P., Hawkins T., Hillier L., Jier M., Johnston L., Jones M., Kershaw J., Kirstein J., Laird N., Latreille P., Lightning J., Lloyd C., McMurray A., Mortimore B., O'Callaghan M., Parsons J., Percy C., Rifkin L., Roopra A., Saunders D., Shownkeen R., Thivierge-Mieg J., Thomas K., Vaughan K., Waterston R., Watson A., Weinstock L., Wilkinson Sproat J., Wohldman P.; elegans.";							
RT	2.2 Mb of contiguous nucleotide sequence from chromosome III of C. elegans.							
RL	Nature 368:32-38(1994).							
RN	[2]	SEQUENCE FROM N.A.						
RC	SEQUENCE=BRISTOL N2;							
RA	Blanchard M., Bradshaw H.; Submitted (OCT-1996) to the EMBL/GenBank/DDBJ databases.							
RN	[3]	SEQUENCE FROM N.A.						
RP	SEQUENCE=BRISTOL N2;							
RC	STRAIN=BRISTOL N2;							
RA	Waterson R;							
RL	Submitted (SEP-1996) to the EMBL/GenBank/DDBJ databases.							
CC	-1- SIMILARITY: TO PANCREATIC TRYPSIN INHIBITOR (KUNITZ) DOMAIN.							
DR	EMBL: U70857; AAB9170.1; -.							
DR	InterPro: IPR002223; Kunitz_BPTI.							
DR	Pfam: PF0014; Kunitz_BPTI; 1.							
DR	SMART: SMD0013; RU; 1.							
DR	PROSITE: PS00280; BPTI_KUNITZ_1; 1.							
DR	PROSITE: PS00279; BPTI_KUNITZ_2; 1.							
KW	Serine protease inhibitor.							
SQ	SEQUENCE 208 AA; 24008 MW; 779ABA8948E67B0 CRC64;							
RESULT	7							
Q9BDM2	Q9BDM2	PRELIMINARY;	PRT;	322 AA.				
ID	Q9BDM2;							
AC	Q9BDM2;							
DT	01-JUN-2001 (TREMBLrel. 17, Created)							
DT	01-JUN-2001 (TREMBLrel. 17, Last sequence update)							
DT	01-JUN-2001 (TREMBLrel. 17, Last annotation update)							
DE	UNKNOWN (PROTEIN FOR IMAGE 619669) (FRAGMENT).							
OS	Homo sapiens (Human).							
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.							
OC	NCBI_TAXID=96006;							
RN	[1]	SEQUENCE FROM N.A.						
RP	TISSUE=LYMPHOMA;							
RC	STRAUBERG R;							
RA	Submitted (MAR-2001) to the EMBL/GenBank/DDBJ databases.							
RL	EMBL: BC004950; AAH04950.1; -.							
DR	NON_TER_1							
FT	SEQUENCE 322 AA; 34916 MW; 33C65BB8B8F9D761 CRC64;							
SQ								
RESULT	8							
Q9DR6	Q9DR6	PRELIMINARY;	PRT;	409 AA.				
ID	Q9DR6;							
AC	Q9DR6;							
DT	01-JUN-2001 (TREMBLrel. 17, Created)							
DT	01-JUN-2001 (TREMBLrel. 17, Last sequence update)							
DE	493439B8RIK PROTEIN.							
GN	493439B8RIK.							
OS	Mus musculus (Mouse).							
OC	Eukaryota; Metazoa; Rodentia; Sciurognathii; Muridae; Murinae; Mus.							
OC	Mammalia; Eutheria; Rodentia; Sciurognathii; Muridae; Murinae; Mus.							
OC	NCBI_TAXID=10090;							
RN	[1]	SEQUENCE FROM N.A.						
RP	SEQUENCE FROM N.A.							
RC	STRAIN=BRISTOL N2;							

STRAIN=C57BL/6J; TISSUE=TESTIS;
MEDLINE=21088660; PubMed=11217851;

RC Kawai J., Shinagawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
RX RA Arakawa T., Kura A., Fukunishi Y., Konno H., Adachi J., Fukuda S.,
RA Kondo S., Yamamoto I.,
RA Aizawa T., Izawa M., Kiyoasa K.,
RA Saito T., Okazaki Y., Gojohori T., Bono H., Kasukawa T., Saito R.,
RA Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,
RA Fleischmann W., Gaasterland T., Gissi C., King B., Kochiwa H.,
RA Kuehl P., Lewis S., Matsuo Y., Nikaido I., Pesole G., Quackenbush J.,
RA Schriml L.M., Staubli F., Suzuki R., Tomita M., Wagner L., Washio T.,
RA Salari K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,
RA Blaize J., Okido T., Bojunga N., Carninci P., de Bonaldo M.F.,
RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Garibaldi M.,
RA Gustincich S., Hill D., Hofmann M., Hume D.A., Kamiy M., Lee N.H.,
RA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Montaerts P.,
RA Norzine P., Ring B., Rodriguez I., Satoh Y., Shibata Y., Storch K.-F.,
RA Sasaki H., Sato K., Schoenbach C., Seya T.,
RA Suzuki H., Toyo-oka K., Wang K.H., Weitz C., Whittaker C., Wilming L.,
RA Wyshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohtsuki S.,
RA Hayashizaki Y.;
RT "Functional annotation of a full-length mouse cDNA collection.";
RL Nucleic Acids Res. 2001; 29:695-696.
DR EMBL; AK017114; BAB3064.1; -.
DR MGD; MGI:19184456; 4933439B08Rik.
DR InterPro; IPR003953; AAA.
DR Pfam; PF00000; AAA; 1.
DR SMART; SM00322; AAA; 1.
DR PROSITE; PS00463; Zn2_CY6_fungal.
SQ SEQUENCE 409 AA; 46.31 MW; 9600B20008DC9749_CRC64;

Query Match 73.9%; Score 34; DB 11; Length 409;
Best Local Similarity 83.3%; Pred. No. 35; Indels 0; Gaps 0;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 2 KLCHQK 7
Db 122 KICHQK 127

RESULT 9

Q9H8F4 ID Q9H8F4 PRELIMINARY; PRT; 522 AA.
AC 09H8F4;
AC 01-WAR-2001 (TREMBLrel. 16, Created)
AC 01-JUN-2001 (TREMBLrel. 16, Last sequence update)
CDNA FLJ13679 ETS, CLONE PLACE2000006.

OS Homo sapiens (Human).
OEUkaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
NCBI_TaxID=9606;

RN [1] RP SEQUENCE FROM N.A.
RC TISSUE=PLACENTA;
RA Isogai T., Ota T., Hayashi K., Sugiyama T., Otsuki T., Suzuki Y.,
RA Nishikawa T., Nagai K., Sugano S., Takahashi-Fujii A., Hara H.,
RA Tanase T., Nomura Y., Togoya S., Komai F., Hara R., Takeuchi K.,
RA Arita M., Nabekura T., Ishii S., Kawai Y., Saito K., Yamamoto J.,
RA Wakamatsu A., Nakamura Y., Nagatari K., Masuho Y., Oshima A.;
RT "NEO human cDNA sequencing project." to the EMBL/GenBank/DBJ databases.
EMBL; AK023741; BAB14633.1; -.
DR InterPro; IPR001849; PH.
DR Pfam; PF001619; PH; 1.
DR SMART; SM006221; RhogEF; 1.
DR SMART; SM00233; PH; 1.
DR PROSITE; PS50003; PH_DOMAIN; 1.
SQ SEQUENCE 522 AA; 59.22 MW; F01A6BF7023920F9_CRC64;

Query Match 73.9%; Score 34; DB 4; Length 522;
Best Local Similarity 62.5%; Pred. No. 42; Indels 0; Gaps 0;
Matches 5; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QKLCHQKK 8
Db 157 QELCHQQE 164

RESULT 10

ID 074308 PRELIMINARY; PRT; 547 AA.
AC 074308;
AC DT 01-NOV-1998 (TREMBLrel. 08, Created)
AC DT 01-JUN-1998 (TREMBLrel. 08, Last sequence update)
AC DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
AC SPBC15D4.02.

OS Schizosaccharomyces pombe (Fission yeast).
OC Schizosaccharomyces pombe; Ascomycota; Schizosaccharomycetes;
OC Schizosaccharomyces; Schizosaccharomyctetes; Schizosaccharomyctaceae;
OC Schizosaccharomyces.
NCBI_TaxID=4896;

RN [1] RP SEQUENCE FROM N.A.
RC STRAIN=972B+;
RA Lyne M., Rajandream M.A., Barrell B.G., Lucas M., Gaillardin C.;
RA Submitted (AUG-1997) to the EMBL/GenBank/DBJ databases.
CC -!- SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY).
CC DOMAIN: EMBL; AL031349; CAA20477.1; -.
DR InterPro; IPR001138; Zn2_CY6_fungal.
DR HSSP; P25502; 1ZME.
DR SMART; SM00066; GAL4; 1.
DR PROSITE; PS00463; ZN2_CY6_FUNGAL_1; 1.
DR PROSITE; PS50048; ZN2_CY6_FUNGAL_2; 1.
KW Zinc_finger; Metal-binding; Nuclear protein; Transcription regulation;
SQ SEQUENCE 547 AA; 59641 MW; A655FD7D039B6CD18_CRC64;

Query Match 73.9%; Score 34; DB 3; Length 547;
Best Local Similarity 71.4%; Pred. No. 43; Indels 0; Gaps 0;

Qy 1 QKLCHQK 7
Db 427 RKLCHER 433

RESULT 11

Q9NVK9 ID Q9NVK9 PRELIMINARY; PRT; 790 AA.
AC 09NVK9;
AC DT 01-OCT-2000 (TREMBLrel. 15, Created)
AC DT 01-OCT-2000 (TREMBLrel. 15, Last sequence update)
CDNA FLJ10665 ETS, CLONE NT2R2005200.

OS Homo sapiens (Human).
OEUkaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
NCBI_TaxID=9606;

RN [1] RP SEQUENCE FROM N.A.
RC TISSUE=PLACENTA;
RA Isogai T., Ota T., Hayashi K., Sugiyama T., Otsuki T., Suzuki Y.,
RA Nishikawa T., Nagai K., Sugano S., Takahashi-Fujii A., Hara H.,
RA Tanase T., Nomura Y., Togoya S., Komai F., Hara R., Takeuchi K.,
RA Arita M., Nabekura T., Ishii S., Kawai Y., Saito K., Yamamoto J.,
RA Wakamatsu A., Nakamura Y., Nagatari K., Masuho Y., Oshima A.;
RT "NEO human cDNA sequencing project." to the EMBL/GenBank/DBJ databases.
EMBL; AK023741; BAB14633.1; -.
DR InterPro; IPR001849; PH.
DR Pfam; PF001619; PH; 1.
DR SMART; SM006221; RhogEF; 1.
DR SMART; SM00233; PH; 1.
DR PROSITE; PS50003; PH_DOMAIN; 1.
SQ SEQUENCE 522 AA; 59.22 MW; F01A6BF7023920F9_CRC64;

DR	EMBL: AK001527; BAA91741.1;	-	CC	-!- SIMILARITY: TO RNA-DIRECTED DNA POLYMERASE (REVERSE TRANSCRIPTASE);	
DR	InterPro; IPR001849; PH.	-	CC	EMBL: AF018033; AAB71689.1;	
DR	InterPro; IPR002019; RhogEF.	-	DR	InterPro; IPR002156; RNaseH.	
DR	Pfam; PF00169; PH; 1.	-	DR	InterPro; IPR000477; RVTse.	
DR	Pfam; PF00621; RhogEF; 1.	-	DR	Pfam; PF00075; rnaseH; 1.	
DR	SMART; SM00233; PH; 1.	-	DR	Pfam; PF00078; rvt; 1.	
DR	SMART; SM00325; RhogEF; 1.	-	KW	RNA-directed DNA Polymerase	
DR	PROSITE; PS50003; PH DOMAIN; 1.	-	SEQUENCE	1295 AA; 144307 MW; E811059B750D5421 CRC64;	
SQ	SEQUENCE 790 AA; 88975 MW; A359F4972CE2DD509 CRC64;	-			
Query Match	Score 34; DB 4; Length 790;	Score 34; DB 3; Length 1295;	Query Match	Score 73.9%; Best Local Similarity 62.5%; Pred. No. 57; Matches 5; Conservative 3; Mismatches 0; Indels 0; Gaps 0;	Score 73.9%; Best Local Similarity 71.4%; Pred. No. 81; Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
Qy	1 QKLCHQK 8	2 KIUCHQK 8	Qy	1 QKLCHQK 8	Score 34; DB 3; Length 1295;
Db	157 QELCHQK 164	33 ELCHQRK 39	Db	33 ELCHQRK 39	Score 73.9%; Best Local Similarity 71.4%; Pred. No. 81; Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
RESULT	12	RESULT	14	RESULT	14
096143	PRELIMINARY; PRT; 1188 AA.	096143	PRELIMINARY; PRT; 1473 AA.	096143	PRELIMINARY; PRT; 1473 AA.
ID	096143	ID	Q63625	ID	Q63625
AC	096143;	AC	Q63625;	AC	Q63625;
DT	01-MAY-1999 (TREMBLrel. 10, Created)	DT	01-NOV-1996 (TREMBLrel. 01, Created)	DT	01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT	01-MAY-1999 (TREMBLrel. 10, Last sequence update)	DT	01-NOV-1998 (TREMBLrel. 01, Last annotation update)	DT	01-NOV-1998 (TREMBLrel. 06, Last annotation update)
DE	PROTEIN WITH 5'-3' EXONUCLEASE DOMAIN (KEM-1 FAMILY).	DE	CBD-BINDING SR-LIKE PROTEIN RA9.	DE	Rattus norvegicus (Rat).
GN	PPB0205C.	GN		OS	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Etheria; Rodentia; Sciurognathi; Muridae; Rattus.
OS	Plasmodium falciparum.	OS		OC	NCBITaxonID=10116;
OC	Eukaryota; Alveolata; Apicomplexa; Haemosporida; Plasmodium.	OC		OX	
OX	NCBI_TaxID=5833;	OX		RN	
RN	[1]	RN		SEQUENCE FROM N.A.	
RP	SEQUENCE FROM N.A.	RP		SEQUENCE FROM N.A.	
RX	MEDLINE=98021743; PubMed=9804551;	RX		TISSUE-HIPPOCAMPUS;	
RA	Gardner M.J., Tettelin H., Carucci D.J., Cummings L.M., Aravind L., Koontz E.V., Shallow S., Mason T., Fujii C., Pederson J., Shen K., Jing J., Aston C., Lai Z., Schwartz D.C., Partea M., Salzberg S., Zhou L., Sutton G.G., Clayton R., White O., Smith H.O., Fraser C.M., Adams M.D., Venter J.C., Hoffman S.L.; RT	RA		MEDLINE=9623459; PubMed=8692929;	
RA	Falciaparum"; Adams M.D., Venter J.C., Hoffman S.L.; RT	RA		Yuryev A., Patterson M., Littingtung Y., Joshi R.V., Gentile C., Gebara M., Corden J.L.; RT	
RA	Science 292:1126-1132(1998).	RA		"The C-terminal domain of the largest subunit of RNA Polymerase II interacts with a novel set of serine/arginine-rich proteins." Proc. Natl. Acad. Sci. U.S.A. 93:6975-6980(1996).	
RL	Exonuclease.	RL		RL	Proc. Natl. Acad. Sci. U.S.A. 93:6975-6980(1996).
DR	AE001380; AAC71830.1; -.	DR		DR	EMBL: U49057; AAC52656.1; -.
SQ	SEQUENCE 1188 AA; 142895 MW;	SQ		SQ	SEQUENCE 1473 AA; 161203 MW; 949BE6F5873989BF CRC64;
Query Match	Score 34; DB 5; Length 1188;	Score 34; DB 11; Length 1473;	Query Match	Score 73.9%; Best Local Similarity 71.4%; Pred. No. 89; Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;	Score 73.9%; Best Local Similarity 71.4%; Pred. No. 89; Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
Qy	1 QKLCHQK 7	1 QKICHQK 7	Qy	1 QKLCHQK 7	Score 34; DB 11; Length 1473;
Db	1117 EELCHAK 1123	Db	1425 QKICHSK 1431	Db	1425 QKICHSK 1431
RESULT	13	RESULT	15	RESULT	15
013348	PRELIMINARY; PRT; 1295 AA.	013348	PRELIMINARY; PRT; 1654 AA.	013348	PRELIMINARY; PRT; 1654 AA.
ID	013348;	ID	Q9PY6	ID	Q9PY6
AC	01-JAN-1998 (TREMBLrel. 05, Created)	AC	Q9PY6;	AC	Q9PY6;
DT	01-JAN-1998 (TREMBLrel. 05, Last sequence update)	DT	01-OCT-2000 (TREMBLrel. 15, Created)	DT	01-OCT-2000 (TREMBLrel. 15, Last sequence update)
DT	01-JUN-2001 (TREMBLrel. 17, Last annotation update)	DT	01-JUN-2001 (TREMBLrel. 17, Last annotation update)	DT	KIAA1542. PROTEIN (FRAGMENT).
DE	REVERSE TRANSCRIPTASE.	DE		GN	Homo sapiens (Human).
OS	Magnaporthe grisea (Rice blast fungus) (Pyricularia grisea).	OS		OS	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Etheria; Primates; Catarhini; Hominidae; Homo.
OC	Scordariomycetes incertae sedis; Magnaportheaceae; Magnaporthe.	OC		OC	NCBITaxonID=9606;
OX	Meyn III M.A., Farrell L., Valente B., Chumley F.G., Orbach M.J.; Submitted (AUG-1997) to the ENSEMBL/GenBank/DDBJ databases.	OX		RN	
RN	[1]	RN		RP	SEQUENCE FROM N.A.
RP	SEQUENCE FROM N.A.	RP		RX	MEDLINE=2077482; PubMed=1081931;
RA	NAgase T., Kikuno R., Ishikawa K., Hirosewa M., Ohara O.; "Prediction of the coding sequences of unidentified human genes.XVII.The complete sequences of 100 new cDNA clones from brain	RA		RA	NAgase T., Kikuno R., Ishikawa K., Hirosewa M., Ohara O.; "Prediction of the coding sequences of unidentified human genes.XVII.The complete sequences of 100 new cDNA clones from brain
RL		RL		RT	

RT which code for large proteins in vitro.";
RL DNA Res. 7:143-150(2000).
CC -I SIMILARITY: CONTAINS A RING-TYPE ZINC FINGER.
EMBL: AB040975; BAA96066.1;
DR InterPro; IPR002219; DAG_Pe-bind.
DR InterPro; IPR001965; PHD.
DR InterPro; IPR001841; Znf_f-ring.
DR Pfam; PF00638; PHD; 1.
DR SMART; SM00109; C1; 1.
DR SMART; SM00249; PHD; 1.
DR PROSITE; PS00184; RING; 2.
DR PROSITE; PS00518; ZINC_FINGER_C3HC4; 1.
KW Zinc-finger.
FT NON_TER 1
SEQUENCE 1654 AA; 179053 MW; 125CF71A84AFFB218 CRC64;
SQ

Query Match 73.9%; Score 34; DB 4; Length 1654;
Best Local Similarity 71.4%; Pred. No. 97;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY	1	QKICHQK	7
		: :	
Db	1606	QKICHSK	1612

Search completed: February 12, 2002, 12:03:26
Job time: 817 sec

Scoring table:	BLOSUM62					
Gapop:	10.0 , Gapext: 0.5					
Searched:	100059 seqs, 366648227 residues					
Total number of hits satisfying chosen parameters:	100059					
Minimum DB seq length:	0					
Maximum DB seq length:	2000000000					
Post-processing:	Minimum Match 0% Maximum Match 100%					
Database :	SwissProt_39:*					
Pred.	No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.					
SUMMARIES						
Result No.	Score	Query	Match	Length	DB ID	Description
1	46	100.0	295	1	BIEA_RAT	P4644 rattus norv
2	36	78.3	654	1	CBP1_YEAST	P07252 saccharomyces
3	33	71.7	673	1	2145_HUMAN	P05516 homo sapien
4	32	69.6	135	1	VE6_APVE	P1131 european el
5	32	69.6	210	1	YQ53_CAEEL	P0311 caenorhabdi
6	32	69.6	272	1	3DHO_ACICA	Q59087 caenorhabd
7	32	69.6	345	1	TF2B_YEAST	P29055 saccharomy
8	32	69.6	357	1	Y303_MYCGE	P47545 mycoplasma
9	32	69.6	595	1	PRIM_CHLTR	Q8A799 chlamydia t
10	32	69.6	609	1	FETA_GORGO	P28050 gorilla gor
11	32	69.6	609	1	FETA_HUMAN	P02771 homo sapien
12	32	69.6	645	1	Y081_CAEEL	P34617 caenorhabdi
13	32	69.6	895	1	SYI_NYCGE	P47587 mycoplasma
14	31	67.4	75	1	EX7S_BACUS	P51522 bacillus su
15	31	67.4	169	1	NEUT_BOVIN	P01156 bos taurus
16	31	67.4	170	1	NEUT_CANFA	P10673 canis famil
17	31	67.4	271	1	YSM4_CAEEL	P01024 caenorhabdi
18	31	67.4	478	1	ARDE_CHLTR	Q84375 chlamydia t
19	31	67.4	498	1	NFS1_SCHPO	Q74351 schizosacch
20	31	67.4	677	1	NRGL_XENLA	P03383 xenopus lae
21	31	67.4	892	1	LDL2_XENLA	Q99088 xenopus lae
22	31	67.4	1207	1	DPOL_ASFB7	P42419 african swi
23	31	67.4	1244	1	DPOL_ASEL6	P42139 african swi
24	31	67.4	1256	1	FLII_DROME	Q24020 drosophila
25	31	67.4	1976	1	MYHA_BOVIN	Q27991 bos taurus
26	31	67.4	1976	1	MYHA_HUMAN	P35580 homo sapien
27	31	67.4	1976	1	MYHA_RAT	Q9j10 rattus norv
28	30	65.2	80	1	EX7S_BACHY	P09968 bacillus ha
29	30	65.2	159	1	MPAA_CORAY	Q88407 corvulus ave
30	30	65.2	259	1	DEC_ECOLI	P00882 escherichia
31	30	65.2	347	1	P75194_MYCPN	P75194 mycoplasma
32	30	65.2	397	1	CAPB_BACAN	P19580 bacillus an
33	30	65.2	442	1	GAG_VILV	P03352 visna lenti
						P23424 visna lenti
						P23425 visna lenti
						P35935 visna lenti
						P16900 ovine lenti
						Q01767 streptomyce
						Q28793 porous tr
						Q57428 xenopus lae
						Q10715 haemobdil
						P56410 anas platyrhynchos
						P02789 gallus gallus
						Q09219 canorhachidi
						Q9wuh7 mus musculus

ALIGNMENTS

RESULT	1
BIEA_RAT	STANDARD;
ID BIEA_RAT	PRT;
AC P4644;	295 AA.
DT 01-NOV-1995 (Rel. 32, Created)	
DT 01-NOV-1995 (Rel. 32, Last sequence update)	
DT 15-JUL-1999 (Rel. 38, Last annotation update)	
DE BILLIVERDIN REDUCTASE A PRECURSOR (EC 1.3.1.24)	(BILLIVERDIN-IX ALPHABETIDASE)
GN BLVR OR BLVR.	
OS Rattus norvegicus (Rat);	
ECO Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Butheria; Rodentia; Murinae; Rattus.	
NCBI TAXID=10116;	
[1]	SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.
TISSUE-Kidney;	
MEDLINE#92156147; PubMed=1371282;	
RA Fakhrai H., Maines M. D.	
RT "Expression and characterization of a cDNA for rat kidney biliverdin reductase. Evidence suggesting the liver and kidney enzymes are the same transcript product."	
RT same transcript product."	
J. Biol. Chem. 267:4023-4029(1992).	
[2]	
RN	
RP MUTAGENESIS IS.	
RX MEDLINE#94291657; PubMed=8020496;	
RA McCoubrey W. K. Jr., Maines M. D.;	
RT site-directed mutagenesis of cysteine residues in biliverdin reductase. Roles in substrate and cofactor binding.	
RL Eur. J. Biochem. 222:1597-603 (1994).	
CC PH OPTIMA USING A DIFFERENT COFACTOR AT EACH PH: NADH AT THE LOWER PH 6.7-6.9 RANGE AND NADPH AT PH 8.5-8.7. NADH, HOWEVER, IS THE PROBABLE COFACTOR IN BIOLOGICAL SYSTEMS.	
CC -!- CATALYTIC ACTIVITY: BILIRUBIN + NAD(P)(+)= BILLIVERDIN + NAD(P)H.	
CC -!- COFACTOR: BILIRUBIN + ONE ZINC ION.	
CC -!- PATHWAY: FINAL STEP IN HEME METABOLISM.	
CC -!- SUBUNIT: MONOMER (BY SIMILARITY).	
CC -!- SUBCELLULAR LOCATION: CYTOPLASMIC.	
CC -!- SIMILARITY: TO E.COLI WHXK.	
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CC EMBL: M81681; AAA40830.1; -	
DR InterPro: IPR000683; GFO_IDH_MoCA.	
DR PFAM: PF01408; GFO_IDH_MoCA; 1.	
KW Oxidoreductase; NAD; NADP; Zinc.	
FT PROPEP	1 2
FT CHAIN	3 295
FT DOMAIN	11 16
	POLY-YVAL.

FT	METAL	279	ZINC (POTENTIAL).	DR SGD; S0003745; CBP1.
FT	METAL	280	ZINC (POTENTIAL).	KW mRNA processing.
FT	METAL	291	ZINC (POTENTIAL).	SO SEQUENCE 654 AA; 76171 MW;
FT	MUTAGEN	292	ZINC (POTENTIAL).	2453B0328EIC44D CRC64;
FT	MUTAGEN	73	C>A: LOSS OF ACTIVITY.	
FT	MUTAGEN	280	C>A: REDUCED ACTIVITY.	Query Match Score 36; DB 1; Length 654;
FT	MUTAGEN	291	C>A: REDUCED ACTIVITY.	Best Local Similarity 75.0%; Pred. No. 21;
SQ	SEQUENCE	295 AA;	239C8EA96C150388 CRC64;	Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
QY	1 QKLCHQKK 8			
Db	288 QKLCHQKK 295			
RESULT 3				
	Z145_HUMAN		STANDARD;	PRT; 673 AA.
ID	Z145_HUMAN			
AC	Q05316;			
DT	01-NOV-1995 (Rel. 32, Created)			
DT	01-NOV-1995 (Rel. 32, Last sequence update)			
DE	ZINC FINGER PROTEIN PLZF (PROMYELOCYTIC LEUKEMIA ZINC FINGER PROTEIN) (ZINC FINGER PROTEIN 145).			
OS	Homo sapiens (Human)			
GN	ZNF145 OR PLZF			
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.			
OC	NCBI_TAXID=9606;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RX	TISSUE=Heart ventricle;			
RX	MEDLINE=9209216; PubMed=8384553;			
RA	Waxman S., Zeleznik A., Chen S.-J., Tong J.-H., Wang Z.-Y., Chen Z., Brand N.J., Chen A., Chen S.-J., Tong J.-H., Wang Z.-Y.,			
RA	"Fusion between a novel Kruppel-like zinc finger gene and the retinoic acid receptor-alpha locus due to a variant t(11;17) translocation associated with acute promyelocytic leukaemia";			
RL	EMBO J. 12:1161-116 (1993). [2]			
RN	SEQUENCE OF 424-455 FROM N.A.			
RX	MEDLINE=92253074; PubMed=8387545;			
RA	Chen S.-J., Zeleznik A., Tong J.-H., Yu H.-Q., Wang Z.-Y., Derré J., Berger R., Waxman S., Chen Z.,			
RA	"Rearrangements of the retinoic acid receptor alpha and promyelocytic leukemia zinc finger genes resulting from t(11;17) (q23;q21) in a patient with acute promyelocytic leukaemia.";			
RT	RL J. Clin. Invest. 91:2260-2267 (1993). [3]			
RX	X-RAY CRYSTALLOGRAPHY (2.0 ANGSTROMS) OF 7-122.			
RA	Marmorstein R.; "Structure-function studies of the BTB/POZ transcriptional repression domain from the promyelocytic leukemia zinc finger oncoprotein.";			
RA	Cancer Res. 59:5275-5282 (1999).			
RA	Li X., Peng H., Schultz D.C., Lopez-Guisa J.M., Rauscher F.J. III, "FUNCTION: PROBABLE TRANSCRIPTION FACTOR. MAY PLAY A ROLE IN MYELOID MATURATION AND/OR MAINTENANCE OF OTHER DIFFERENTIATED TISSUES			
CC	"- SUBCELLULAR LOCATION: NUCLEAR.			
CC	"- ALTERNATIVE PRODUCTS: 2 ISOFORMS; PLZFA AND PLZFB (SHOWN HERE); ARE PRODUCED BY ALTERNATIVE SPLICING.			
CC	"- TISSUE SPECIFICITY: WITHIN THE HEMATOPOIETIC SYSTEM, PLZF IS EXPRESSED IN BONE MARROW, EARLY MYELOID CELL LINES AND PERIPHERAL BLOOD MONONUCLEAR CELLS. ALSO EXPRESSED IN THE OVARY, AND AT LOWER LEVELS, IN THE KIDNEY AND LUNG.			
CC	"- INDUCTION: BY RETINOIC ACID.			
CC	"- DISEASE: A FORM OF ACUTE PROMYELOCYTIC LEUKEMIA (APL) IS CHARACTERIZED BY A CHROMOSOMAL TRANSLOCATION t(11;17)(q32;q21) WHICH INVOLVES ZNF145 AND RETINOIC ACID RECEPTOR ALPHA (RAR α).			
CC	"- SIMILARITY: BELONGS TO THE KRUEPPEL FAMILY OF C2H2-TYPE ZINC-FINGER PROTEINS.			
CC	"- SIMILARITY: CONTAINS 1 BTB/POZ DOMAIN.			
CC				

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CC MEDLINE=87219878; PubMed=3034730;
 CC RX RA Ahola H.; Bergman P.; Stroem A.C.; Moreno-Lopez J.J.; Pettersson U.;
 CC RT RT "Organization and expression of the transforming region from the European elk Papillomavirus (EEPV).";
 CC RT RT Gene 50:1195-205 (1986).
 CC RL CC -!- FUNCTION: EXHIBIT A STRONG, BUT NON SPECIFIC AFFINITY FOR DOUBLE STRANDED DNA (IN VITRO).
 CC CC -!- SUBCELLULAR LOCATION: NUCLEAR MATRIX-ASSOCIATED.
 CC CC DR EMBL: Z19002; CAA79489.1; -.
 CC DR EMBL: S60093; AAC60390.2; -.
 CC DR PDB: ICS3; 09-AUG-00.
 CC DR TRANSFAC; T02336; -.
 CC DR MIM: 176791; -.
 CC DR InterPro; IPR000210; BTB_P0Z.
 CC DR InterPro; IPR000822; znf_C2H2.
 CC DR Pfam: PF0051; BTB; 1.
 CC DR Pfam: PF00096; zf-C2H2; 9.
 CC DR PRINTS; PRO0048; ZINC FINGER.
 CC DR SMART; SM00225; BTB; 1.
 CC DR PROSITE; PS00097; BTB; 1.
 CC DR PROSITE; PS00028; ZINC_FINGER_C2H2_1; 8.
 CC DR PROSITE; PS50157; ZINC_FINGER_C2H2_2; 9.
 CC KW Transcription Regulation; DNA-binding; Zinc-finger; Metal-binding;
 CC KW Nuclear Protein; Repeat; Chromosomal translocation; Proto-oncogene;
 CC KW Phosphorylation; Alternative splicing; 3D-structure.
 FT DOMAIN 34 96 BTB.
 FT 404 652 ZINC FINGERS.
 FT 404 426 C2H2-TYPE.
 FT 432 454 C2H2-TYPE.
 FT 461 483 C2H2-TYPE.
 FT 490 512 C2H2-TYPE.
 FT 518 540 C2H2-TYPE.
 FT 546 568 C2H2-TYPE.
 FT 574 596 C2H2-TYPE.
 FT 602 624 C2H2-TYPE.
 FT 630 652 C2H2-TYPE.
 FT 394 395 BREAKPOINT FOR TRANSLLOCATION TO FORM PLZF-RAR-ALPHA ONCOGENE.
 FT 76 76 PHOSPHORYLATION (BY PDPK) (POTENTIAL).
 FT 184 184 PHOSPHORYLATION (BY PDPK) (POTENTIAL).
 FT 197 197 PHOSPHORYLATION (BY PDPK) (POTENTIAL).
 FT 256 256 PHOSPHORYLATION (BY PDPK) (POTENTIAL).
 FT 282 282 PHOSPHORYLATION (BY PDPK) (POTENTIAL).
 FT 628 628 PHOSPHORYLATION (BY PDPK) (POTENTIAL).
 FT 255 377 MISSING (IN ISOFORM PLZFA).
 SQ SEQUENCE 673 AA; 74332 MW; 7CD719E2A32109D CRC64;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=BRISTOL N2;
 RA Favello T;
 RL Submitted (JUL-1995) to the EMBL/GenBank/DDBJ databases.
 CC -!- SIMILARITY: IC C_ELEGANS 2K675.1.
 CC
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 CC
 EMBL; U23176; AAC46715.1; -.
 DR WormPep; F21H12.3; CE01914.
 DR Hypothetical Protein.
 KW SEQUENCE 210 AA; 23617 MW; 1E646EEFC30154AO CRC64;
 SQ
 Query Match 71.7%; Score 33; DB 1; Length 673;
 Best Local Similarity 83.3%; Pred. No. 75; Indels 0; Gaps 0;
 Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 Qy 2 KLCHQK 7
 Db 605 KLCHQR 610
 RESULT 4
 V6_PAPV ID_V6_PAPV STANDARD; PRT; 135 AA.
 AC P11331; 11, Created
 DT 01-JUL-1989 (Rel. 11, Last sequence update)
 DT 01-JUL-1989 (Rel. 11, Last annotation update)
 DE EG PROTEIN.
 GN EG
 OS European elk Papillomavirus (EEPV).
 OC Viruses; dsDNA viruses, no RNA stage; Papillomaviridae;
 OC Papillomavirids
 OX NCBI_TaxID=10565;
 RN [1]
 RP SEQUENCE FROM N.A.

DE TRANSCRIPTION INITIATION FACTOR IIB (TFIIB) (TRANSCRIPTION FACTOR E).
 GN SUA7 OR IPR086W OR P9513_4.
 OS Saccharomyces cerevisiae (Baker's yeast)
 OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomyces;
 OC Saccharomycetales; Saccharomyctaceae; Saccharomyces.
 NCBI_TaxID=4932;
 OX
 RN [1] SEQUENCE FROM N.A.
 RP SEQUENCE FROM N.A.
 MEDLINE=92191276; PubMed=1547497;
 RX
 RA Pinto I.; Ware D.E.; Hampsey M.;
 RT "The yeast SUA7 gene encodes a homolog of human transcription factor
 TFIIB and is required for normal start site selection in vivo."
 RT Cell 68:977-988(1992).
 RL [2] SEQUENCE FROM N.A.
 RP STRAINS288C / AB972;
 RC Johnston M., Andrews S., Brinkman R., Cooper J., Ding H., Du Z.,
 RA Favello A., Fulton L., Gattung S., Greco T., Hawkins J., Hillier L., Kirsten J.,
 RA Kucaba T., Hollsworth K., Hawkins J., Hillier L., Tier M.,
 RA Johnson D., Johnston L., Langston Y., Latreille P., Le T.,
 RA Mardis E., Menezes S., Miller N., Shan M., Pauley A., Peluso D.,
 RA Rifkin L., Riles L., Taich A., Treskakis P., Vignetti D.,
 RA Wilcox L., Wohlgemut P., Vaughn M., Wilson R., Waterston R.;
 RL Submitted (MAR-1996) to the EMBL/GenBank/DBJ databases.
 RT FUNCTION: GENERAL FACTOR THAT PLAYS A MAJOR ROLE IN THE ACTIVATION
 CC OF EUKARYOTIC GENES TRANSCRIBED BY RNA POLYMERASE II.
 CC SUBUNIT: ASSOCIATES WITH TFIID-LIA (DA COMPLEX) TO FORM TFIID-LIA.
 CC TFIID-LIA (DAB-COMPLEX) WHICH IS THEN RECOGNIZED BY POLYMERASE II.
 CC !- SUBCELLULAR LOCATION: NUCLEAR
 CC !- SIMILARITY: BELONGS TO THE TFIIB FAMILY.
 CC -----
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 CC or send an email to license@isb-sib.ch).
 CC -----
 DR M81380; AAA35126.1; -.
 DR U51033; AAB68135.1; -.
 DR S26707; S26707.
 DR Q00403; ITTB.
 DR HSSP; T00819; -.
 DR TRANSFAC; T00819; -.
 DR S0006380; SUA7; -.
 DR InterPro; IPR000553; Cyclin.
 DR InterPro; IPR000553; TFIIB.
 DR PFAM; PF00382; transcript_fact2; 2.
 DR PRINTS; PRO00685; TFIIFCTORIIB.
 DR SMART; SM00385; CYCLIN; 2.
 DR PROSITE; PS00782; TFIIB; 1.
 DR Transcription regulation: Nuclear protein; Zinc-finger.
 KW
 FT ZNFING 24 48 POTENTIAL.
 FT REPEAT 136 212
 FT REPEAT 242 318
 SQ SEQUENCE 345 AA; 38200 MW; 8F1FD24602436E2 CRC64;

Query Match 69 6%; Score 32; DB 1; Length 345;
 Best Local Similarity 85.7%; Pred. No. 51; Indels 0; Gaps 0;
 Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 KLCHQKK 8
 DB 149 KLAHQKK 155

RESULT 7
 TF2B_YEAST STANDARD PRT; 345 AA.
 ID TF2B_YEAST
 AC P90555;
 DT 01-DEC-1992 (Rel. 24, Created)
 DT 01-DEC-1992 (Rel. 24, Last sequence update)
 DT 15-JUL-1998 (Rel. 36, Last annotation update)

QY 2 KLCHQKK 8
 DB 155 KUCHEDEK 161

RESULT 8
 Y303_MYCGE STANDARD PRT; 357 AA.
 ID Y313_MVCGE
 AC P41545;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)

DT 20-AUG-2001 (Rel. 40, Last annotation update)
 GN HYPOTHETICAL ABC TRANSPORTER ATP-BINDING PROTEIN MG303.
 OS Mycoplasma genitalium.
 OC Firmicutes; Bacillus/Clostridium group; Mollicutes;
 OC Mycoplasmataceae; Mycoplasma.
 OX NCBI_TaxID=2097;

RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=ATCC 33530 / G-37;
 RX MEDLINE#=96026346; PubMed=7569993;
 RA Fraser C.M., Gocayne J.D., White O., Adams M.D., Clayton R.A.,
 RA Sutton G., Kerlavage A.R.,
 RA Fleischmann R.D., Bult C.J., Ketchum KA., Kelley J.M.,
 RA Fritchman J.L., Weidman J.F., Small K.V., Sandusky M., Fuhrmann J.L.,
 RA Nguyen D.T., Utterback T.R., Sauder D.M., Phillips C.A., Merrick J.M.,
 RA Tomb J.-F., Dougherty B.A., Boit K.F., Hu P.C., Lucier T.S.,
 RA Peterson S.N., Smith H.O., Hutchison C.A. III, Venter J.C.;
 RT "The minimal gene complement of Mycoplasma genitalium."
 RL Science 270:197-203 (1995).
 -!- SIMILARITY: BELONGS TO THE ATP-BINDING TRANSPORT PROTEIN FAMILY
 (ABC TRANSPORTERS).

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CC SEQUENCE FROM N.A.
 RC STRAIN=ATCC 33530 / G-37;
 RX MEDLINE#=96026346; PubMed=7569993;
 RA Fraser C.M., Gocayne J.D., White O., Adams M.D., Clayton R.A.,
 RA Sutton G., Kerlavage A.R.,
 RA Fritchman J.L., Weidman J.F., Small K.V., Sandusky M., Fuhrmann J.L.,
 RA Nguyen D.T., Utterback T.R., Sauder D.M., Phillips C.A., Merrick J.M.,
 RA Tomb J.-F., Dougherty B.A., Boit K.F., Hu P.C., Lucier T.S.,
 RA Peterson S.N., Smith H.O., Hutchison C.A. III, Venter J.C.;
 RT "The minimal gene complement of Mycoplasma genitalium."
 RL Science 270:197-203 (1995).
 -!- SIMILARITY: BELONGS TO THE ATP-BINDING TRANSPORT PROTEIN FAMILY
 (ABC TRANSPORTERS).

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CC SEQUENCE FROM N.A.
 DR TIGR; MG303; IPRO03593; AAA.
 DR InterPro; IPRO03439; ABC_TRANSPORTER.
 DR InterPro; IPRO01687; ATP_GTP_A.
 DR SMART: SM00382; AAA; 1.
 DR PROSITE; PS00211; ABC_TRANSPORTER; FALSE_NEG.
 KW Hypothetical Protein; ATP-binding; Transport; Complete proteome.
 FT NP BIND 107 114 ATP (POTENTIAL)
 SQ SEQUENCE 357 AA; 40786 MW; AFB102F88E090E CRC64;

Query Match 69.6% Score 32; DB 1; Length 357;
 Best Local Similarity 83.3%; Pred. No. 65;
 Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy' 2 KLCHQ 7
 Db 1.83 KLCHKK 188

RESULT 9
 PRIM_CHLTR STANDARD; PRT; 595 AA.
 ID PRIM_CHLTR
 AC 084799;
 DT 20-AUG-2001 (Rel. 40, Created)
 DT 20-AUG-2001 (Rel. 40, Last sequence update)
 DE DNA PRIMASE (EC 2.7.7.-).
 GN DNAG OR CTT94.
 OS Chlamydia trachomatis.
 OC Bacteria; Chlamydiales; Chlamydiaceae; Chlamydia.
 OX NCBI_TaxID=813;
 RN SEQUENCE FROM N.A.
 RC STRAIN=D/UV/3/CX;
 RX MEDLINE#=99000800; PubMed=9784136;
 RA Stephens R.S.; Kalman S.; Lammel C.J.; Fan J.; Marathe R.; Aravind L.;
 RA Mitchell W.P.; Olinger L.; Tatusov R.L.; Zhao Q.; Koornin E.V.;
 RA Davis R.W.;
 RT "Genome sequence of an obligate intracellular pathogen of humans: Chlamydia trachomatis.";
 RT

RL Science 282:754-759(1998).
 -!- FUNCTION: DNA PRIMASE IS THE POLYMERASE THAT SYNTHESIZES SMALL RNA PRIMERS FOR THE OKAKAII FRAGMENTS ON BOTH TEMPLATE STRANDS AT REPLICATION FORKS DURING CHROMOSOMAL DNA SYNTHESIS.
 CC -!- COFACTOR: BINDS ONE ION PER MOLECULE (BY SIMILARITY).
 CC -!- SUBUNIT: MONOMER (BY SIMILARITY).
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 CC EMBL: AE001351; AAC68389.1;
 DR InterPro; IPRO02936; Toprism;
 DR InterPro; IPRO06944; ZNF_CHC2C;
 DR Pfam; PF01751; Toprism; 1;
 DR ProDom; PD002988; zf-CHC2; 1;
 DR SMART; SN04040; ZNF_CHC2; 1;
 DR Transferase; DNA replication; DNA-directed RNA polymerase; Primosome;
 KW Zinc; Metal-binding; Complete proteome.
 KW Zinc; Metal-binding; Zinc finger protein.
 FT 2N_FING 38 62
 SQ SEQUENCE 595 AA; 536858EBAFCD8FB6 CRC94;
 Query Match 69.6% Score 32; DB 1; Length 595;
 Best Local Similarity 83.3%; Pred. No. 1e+02;
 Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 QKLOHQ 6
 Db 565 RKLCHQ 570

RESULT 10
 FETA_GORGEO STANDARD; PRT; 609 AA.
 ID FETA_GORGEO
 AC P28050;
 DT 01-AUG-1992 (Rel. 23, Created)
 DT 01-AUG-1992 (Rel. 23, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE ALPHA-FETOPROTEIN PRECURSOR (ALPHA-FETOPROTEIN).
 DE AFP.
 GN Gorilla gorilla gorilla (Lowland gorilla).
 OS Gorilla gorilla; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Gorilla.
 OC NCBL_TaxID=9552;
 OX [1]
 RN SEQUENCE FROM N.A.; PubMed=1706310;
 RP MEDLINE#=1169317;
 RX Ryan S.C., Zielinski R., Dugaiczky A.;
 RT "Structure of the gorilla alpha-fetoprotein gene and the divergence of primates";
 RT Genomics 9:60-72 (1991).
 CC -!- FUNCTION: BINDS COPPER, NICKEL, AND FATTY ACIDS AS WELL AS, AND BILIRUBIN LESS WELL THAN, SERUM ALBUMEN.
 CC -!- SUBUNIT: DIMERIC AND TRIMERIC FORMS HAVE BEEN FOUND IN ADDITION TO THE MONOMERIC FORM (BY SIMILARITY).
 CC -!- SUBCELLULAR LOCATION: EXTRACELLULAR.
 CC -!- TISSUE SPECIFICITY: PLASMA.
 CC -!- DOMAIN: COMPOSED OF THREE HOMOLOGOUS DOMAINS.
 CC -!- PTM: SULFATED (BY SIMILARITY).
 CC -!- SIMILARITY: BELONGS TO THE ALB/APP/VDB FAMILY.

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[1] RN SEQUENCE OF 1-28 FROM N.A.
RN MEDLINE=33278385; PubMed=7684942;
RX
RA McVey J.H., Michaelides K., Hansen L.P., Ferguson-Smith M.,
RA Tilmann S., Krumlau R., Tudman E.G.D.;
RT "A G-->A substitution in an HNF 1 binding site in the human alpha-fetoprotein gene is associated with hereditary persistence of alpha-fetoprotein (HFAFP).";
RT
RN Genet. 2:379-379(1993).
RN
RN SEQUENCE OF 429-556 FROM N.A.
RX MEDLINE=8-158778; PubMed=5187626;
RA Beattie W.G., Dugaiczyk A.;
RA Fucci P., Siciliano R., Malorni A., Marino G., Tecce M.F.,
RA Ceccarini C., Terrana B.;
RT "Structure and evolution of human alpha-fetoprotein deduced from partial sequence of cloned cDNA.";
RT
RN Gene 20:415-422(1982).
RN
RN PARTIAL SEQUENCE OF 19-609.
RX MEDLINE=91242409; PubMed=1709810;
RA
RA Fucci P., Siciliano R., Malorni A., Marino G., Tecce M.F.,
RA Ceccarini C., Terrana B.;
RT "Human alpha-fetoprotein primary structure: a mass spectrometric study";
RT
RN Biochemistry 30:5061-5066(1991).
RN
RN PRELIMINARY SEQUENCE OF 19-35.
RX MEDLINE=77242506; PubMed=0228;
RA Yachnin S., Hsu R., Heinrikson R.L., Miller J.B.;
RT "Studies on human alpha-fetoprotein. Isolation and characterization of monomeric and polymeric forms and amino-terminal sequence analysis.";
RT
RN Biochim. Biophys. Acta 493:418-428(1977).
RN
RN PARTIAL SEQUENCE OF 19-39.
RX MEDLINE=75018719; PubMed=4138095;
RA Rosolanti E., Palko H., Vahteri A., Seppala M., Virolainen M.,
RA Konttinen A.;
RT "Alpha fetoprotein: structure and expression in man and inbred mouse strains under normal conditions and liver injury.";
RT
RN Cancer Res. 37:3663-3667(1977).
RN
RN PRELIMINARY SEQUENCE OF 19-39.
RX MEDLINE=75018719; PubMed=4138095;
RA Rosolanti E., Palko H., Vahteri A., Seppala M., Virolainen M.,
RA Tamaoki T.,
RT "The human alpha-fetoprotein gene. Sequence organization and the 5' flanking region.";
RT
RN J. Biol. Chem. 260:5055-5060(1985).
RN
RN GENE STRUCTURE.
RX MEDLINE=85188629; PubMed=2580830;
RA
RA Aoyagi Y., Ikenaka T., Ichida F., Wegmann T.G.,
RA Tamaoki T.;
RT "The human alpha-fetoprotein gene. Sequence organization and the 5' flanking region.";
RT
RN Cancer Res. 38:3483-3486(1978).
RN
RN METAL-BINDING.
RX MEDLINE=79001617; PubMed=80265;
RA Aoyagi Y., Ikenaka T., Ichida F.;
RT "Copper(II)-binding ability of human alpha-fetoprotein.";
RT
RN Cancer Res. 38:3483-3486(1978).
RN
RN BILIRUBIN-BINDING.
RX MEDLINE=80001710; PubMed=89900;
RA Aoyagi Y., Ikenaka T., Ichida F.;
RT "Alpha-fetoprotein as a carrier protein in plasma and its bilirubin-binding ability.";
RT
RN Cancer Res. 39:3571-3574(1979).
RN
RN SULFATION.
RX PubMed=2414772;
RA Liu M.C., Yu S., Sy J., Redman C.M., Lipmann F.;
RT "Tyrosine sulfation of proteins from the human hepatoma cell line [12]."

RT HepG2. ";
 RT Proc. Natl. Acad. Sci. U.S.A. 82:7160-7164 (1985).
 CC -!- FUNCTION: BINDS COPPER, NICKEL, AND FATTY ACIDS AS WELL AS, AND
 CC -!- BILIRUBIN LESS WELL THAN, SERUM ALBUMIN. ONLY A SMALL PERCENTAGE
 CC -!- (LESS THAN 2%) OF THE HUMAN AFP SHOWS ESTROGEN-BINDING PROPERTIES.
 CC -!- SUBUNIT: DIMERIC AND TRIMERIC FORMS HAVE BEEN FOUND IN ADDITION
 CC -!- TO THE MONOMERIC FORM.
 CC -!- TISSUE SPECIFICITY: PLASMA. SYNTHESIZED BY THE FETAL LIVER AND
 CC -!- YOLK SAC.
 CC -!- DEVELOPMENTAL STAGE: OCCURS IN THE PLASMA OF FETUSES MORE THAN 4
 CC -!- WEEKS OLD, REACHES THE HIGHEST LEVELS DURING THE 12TH-16TH WEEK OF
 CC -!- GESTATION, AND DROPS TO TRACE AMOUNTS AFTER BIRTH. THE SERUM LEVEL
 CC -!- IN ADULTS IS USUALLY LESS THAN 40 NG/ML. AFP OCCURS ALSO AT HIGH
 CC -!- LEVELS IN THE PLASMA AND ASCitic FLUID OF ADULTS WITH HEPATOMA.
 CC -!- DOMAIN: COMPOSED OF THREE HOMOLOGOUS DOMAINS.
 CC -!- PTM: INDEPENDENT STUDIES SUGGEST HETEROGENEITY OF THE AMINO-
 CC -!- TERMINAL SEQUENCE OF THE MATURE PROTEIN AND OF THE CLEAVAGE SITE
 CC -!- OF THE SIGNAL SEQUENCE.
 CC -!- PTM: SULFATED.
 CC -!- SIMILARITY: BELONGS TO THE ALB/AFP/YDB FAMILY.

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 CC or send an email to license@isb-sib.ch).

CC -!- DR: M10349; AAA51674.; -;
 CC -!- DR: M10350; AAA51675.; -;
 CC -!- DR: V01514; CAA24158.; -;
 CC -!- DR: M16110; AAB58154.; -;
 CC -!- DR: EMBL: J00077; AAC95396.; -;
 CC -!- DR: EMBL: Z19532; CAA79592.; -;
 CC -!- DR: PIR: A03234; PPHU.; -;
 CC -!- DR: PIR: A26624; A26624.; -;
 CC -!- DR: HSSP: P02768; I1UOR.
 CC -!- DR: GLYCOSUITEDB; P02771; -;
 CC -!- DR: MIM: 104150; -;
 CC -!- DR: InterPro: IPR000264; Serum_albumin.
 CC -!- DR: Pfam: PF00273; transport_prot; 3.
 CC -!- DR: PRINTS: PRO0802; SERMALBUMIN.
 CC -!- DR: SMART: SM00103; ALBUMIN; 3.
 CC -!- DR: PROSITE: PS00212; ALBUMIN; 2.
 CC -!- KW: Glycoprotein; Sulfation; Albumin; Plasma; Embryo; Repeat;
 CC -!- KW: Metal-binding; Copper; Nickel; Signal.
 CC -!- FT: SIGNAL: 1¹⁸ ALPHA_FETOPROTEIN.
 CC -!- FT: CHAIN: 19 609 COPPER AND NICKEL.
 CC -!- FT: METAL: 22 22 COPPER AND NICKEL.
 CC -!- FT: DISULFID: 99 114
 CC -!- FT: DISULFID: 113 124
 CC -!- FT: DISULFID: 148 193
 CC -!- FT: DISULFID: 192 201
 CC -!- FT: DISULFID: 224 270
 CC -!- FT: DISULFID: 269 277
 CC -!- FT: DISULFID: 289 303
 CC -!- FT: DISULFID: 302 313
 CC -!- FT: DISULFID: 384 393
 CC -!- FT: DISULFID: 416 462
 CC -!- FT: DISULFID: 461 472
 CC -!- FT: DISULFID: 485 501
 CC -!- FT: DISULFID: 500 511
 CC -!- FT: DISULFID: 538 583
 CC -!- FT: DISULFID: 582 591
 CC -!- FT: CARBOHYD: 251 251 N-LINKED (GLCNAC. . .).
 CC -!- SQ: SEQUENCE: 609 AA: 68677 MW: 4D4E5820E1C2D4F CRC64;

Best Local Similarity 50.0%; Pred. No. 1e+02; Indels 0; Gaps 0;
 Matches 4; Conservative 4; Mismatches 0;

Qy 1 QKLHQKK 8
 Db 96 EBLHERE 103

RESULT 12

Y081_CAEEL STANDARD; PRT: 645 AA.

ID P34617;
 AC Y081_CAEEL
 DT 01-FEB-1994 (Rel. 28; Created)
 DT 01-FEB-1995 (Rel. 31; Last sequence update)
 DT 20-AUG-2001 (Rel. 40; Last annotation update)

HYPOTHETICAL GTP-BINDING PROTEIN ZK1236.1 IN CHROMOSOME III.

DE ZK1236.1.
 GN
 OS Ceanorhabditis elegans.
 OC Nematoda; Chromadorea; Rhabditida; Rhabditoidea;
 OC Rhabditidae; Peioderinae; Caenorhabditis;
 NCBI_TAXID=6239;

OX RN

RP SEQUENCE FROM N_A.
 RC STRAIN-BRISTOL N2;
 RX MEDLINE-94150718; PubMed=7906398;

RA Wilson R., Ainscough R., Anderson K., Baynes C., Berks M., Bonfield J., Burron J., Connell M., Copsey T., Cooper J., Coulson A., Craxton M., Dear S., Du Z., Durbin R., Favell A., Fraser A., Fulton L., Gardner A., Green P., Hawkins T., Hillier L., Jier M., Johnston L., Jones M., Kershaw J., Kirsten J., Laird N., Latreille P., Lightning J., Lloyd C., Mortimore B., O'Callaghan M., Parsons J., Percy C., Rifkin L., Roopra A., Saunders D., Showkeen R., Sims M., Smalton N., Smith A., Sonnhammer E., Staden R., Sulston J., Thierry-Mieg J., Thomas K., Vaughan K., Watson R., Watson A., Weinstock L., Wilkinson-Sprout J., Woldman P.;
 RA "2.2 Mb of contiguous nucleotide sequence from chromosome III of C. elegans." RT Nature 368:32-38(1994).
 RL "-!- SIMILARITY: BELONGS TO THE GTP-BINDING ELONGATION FACTOR FAMILY.

CC LEPA SUBFAMILY.

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CC -!- DR: EMBL: 113200; AAA28191.; -;
 CC -!- DR: HSSP: P13551; IEL0.
 CC -!- DR: WormPeP: ZK1236.1; CEO1446.
 CC -!- DR: InterPro: IPR000075; GTP_EFTU.
 CC -!- DR: Pfam: PF00009; GTP_EFTU.; 1.
 CC -!- DR: PROSITE: PS00001; EFACTOR_GNP; 1.
 CC -!- KW: Hypothetical protein; GTP-Binding.
 CC -!- FT: NP_BIND: 49 56 GTP (POTENTIAL).
 CC -!- FT: NP_BIND: 108 112 GTP (POTENTIAL).
 CC -!- FT: NP_BIND: 162 165 GTP (POTENTIAL).
 CC -!- SQ: SEQUENCE 645 AA: 72268 MW: 3F08E3E5FD53819_CRC64;

Query Match Score 32; DB 1; Length 645;
 Best Local Similarity 75.0%; Pred. No. 1.1e+02;
 Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 QKLHQKK 8
 Db 614 KKLSHQKK 621

RESULT 13

RP SEQUENCE FROM N.A.
MEDLINE-8819625; PubMed=2832414;
RX Kislauskis E., Bullock B., McNeil S., Dobner P.R.;
RT "The rat gene encoding neurotensin and neuropeptide N. Structure,
RT tissue-specific expression, and evolution of exon sequences.";
RL J. Biol. Chem. 263:4963-4968 (1988).
RN [2]
RP SEQUENCE OF 150-162.
RT TISSUE-Hypothalamus;
RX MEDLINE-75095679; PubMed=1112838;
RA Carraway R., Leeman S.E.;
RA Carraway R., Leeman S.E.;
RT "The amino acid sequence of a hypothalamic peptide, neuropeptins.";
RT J. Biol. Chem. 250:1907-1911(1975).
RN [3]
RN SYNTHESIS OF NEUROTENSIN.
RX MEDLINE-75095679; PubMed=1112838;
RA Carraway R., Leeman S.E.;
RA Carraway R., Leeman S.E.;
RT "The synthesis of neurotensin.";
RT J. Biol. Chem. 250:1912-1918(1975).
CC -I- FUNCTION: NEUROTENSIN MAY PLAY AN ENDOCRINE OR PARACRINE ROLE
CC IN THE REGULATION OF FAT METABOLISM. IT CAUSES CONTRACTION OF
CC SMOOTH MUSCLE.
CC -I- SUBCELLULAR LOCATION: PACKAGED WITHIN SECRETORY VESICLES.
CC -I- TISSUE SPECIFICITY: BRAIN AND GUT.
CC -I- SIMILARITY: C-TERMINAL SEQUENCE SIMILARITY WITH NEUROTENSIN-
CC RELATED PEPTIDES.
CC -----
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CC -----
CC EMBL, M18621; AAA30668.1; -.
DR PIR; A01420; UNBO.
DR Cleavage on pair of basic residues; Vasoactive; Signal.
KW SIGNAL 1 22 POTENTIAL.
FT SIGNAL 1 22 POTENTIAL.
FT CHAIN 23 147 LARGE NEUROMEDIN N (NMN-125).
FT PEPTIDE 142 147 NEUROMEDIN N.
FT PEPTIDE 150 162 NEUROTENSIN.
FT MOD_RES 150 150 PYRROLIDINE CARBOXYLIC ACID.
FT SO SEQUENCE 169 AA: 19712 MW: 7B78100D5E9D32 CRO64;
FT MOD_RES 169 AA: 19712 MW: 7B78100D5E9D32 CRO64;
FT SO SEQUENCE 169 AA: 19712 MW: 7B78100D5E9D32 CRO64;

```

Query Match 67.4%; Score 31; DB 1; Length 169;
Best Local Similarity 57.1%; Pred. No. 51;
Matches 4; Conservative 2; Mismatches 1; Indels 0;
Gaps 0;

```

QY 1 QKLCHQK 7
DB 107 QKLCHQD 112

Search completed: February 12, 2002, 12:04:02
Job time: 798 sec

Copyright	GenCore version 4.5 (c) 1993 - 2000 Compugen Ltd.	
protein - protein search, using sw model	February 12, 2002, 11:51:40 ; Search time 55.4 Seconds (without alignments) 11.000 Million cell updates/sec	
title:	US-09-606-129A-19	
perfect score:	46	
sequence:	1 QKLCHOKK 8	
scoring table:	BLOSUM62	

total number of hits containing chosen connectors.

length seq DB minimum DB

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Post-processing: Minimum Match 0%

Maximilian March 1903
Listings first 45 summaries

PIR_681*
1 2251*

```
2:    pir2:*
```

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No. is the number of results predicted by chance to have a score greater than or equal to the score of the total score distribution.

SUMMARIES					
Result No.	Score	Query Match	Length DB	ID	Deep
1	46	100.0	295	A42268	bill
2	36	78.3	654	BVBPI1	CBW
3	34	73.9	547	T39478	zir
4	34	73.9	1188	A71621	pro
5	34	73.9	1295	T30282	rea
6	34	73.9	1473	T31422	C-t
7	33	71.7	263	E7275	hyp
8	33	71.7	673	S36336	pre
9	33	71.7	773	T00554	hyp
10	33	71.7	1247	T18571	hyp
11	32	69.6	135	W6WLP	E6
12	32	69.6	210	T16125	hyp
13	32	69.6	290	I39122	3-c
14	32	69.6	345	S26107	tra
15	32	69.6	357	E64333	mem
16	32	69.6	380	T24081	hyp
17	32	69.6	422	E9673	hy
18	32	69.6	462	A46110	te
19	32	69.6	465	T27032	hyp
20	32	69.6	474	T00599	hy
21	32	69.6	548	T22137	hy
22	32	69.6	581	S44816	2K
23	32	69.6	595	F71471	pre
24	32	69.6	609	FPHU	all
25	32	69.6	609	FPGO	all
26	32	69.6	695	B64236	is
27	32	69.6	920	JCT313	ary
28	32	69.6	1056	T00000	2
29	32	69.6	1090	W66666	6.4
30	32	69.6	1100	W66666	2.1

RESULT 1
 A42268
 Biliverdin reductase (EC 1.3.1.24) - rat
 C;Species: Rattus norvegicus (Norway rat)
 C;Date: 04-Mar-1993 #sequence_revision 18-Nov-1
 C;Accession: A42268
 Mairnes, M.D.
 P;Author: P.Khatri, H.;
 J.Biol.Chem. 267, 4023-4029, 1992
 A;Title: Expression and characterization of a c
 A;Reference number: A42268; PMID: 92156147
 A;Accession: A42268
 A;Status: preliminary; not compared with concep
 A;Molecular type: nucleic acid; protein
 A;Residues: 1-295 <PAK>
 A;Cross-references: GB:MB1681; NID:9203177; PID:
 A;Experimental sources: kidney
 A;Sequence extracted from NCBI backbone (<
 C;Keywords: liver; oxidoreductase

```

Query Match          100.0% Score 46; D
Best Local Similarity 100.0% Pred. No. 0.
Matches           8; Conservative 0; Mismatches
                    0

Qy      1 QKLCHQK 8
        ||||| |
Db     288 QKLCHQK 295

RESULT          2
BvBYBP1
CBP1 protein - yeast (Saccharomyces cerevisiae)
N; Alternative names: protein HRA54; protein J02;
C; Species: Saccharomyces cerevisiae
C; Date: 31-Mar-1991 #sequence_revision 31-Mar-1991
C; Accession: S03829; S50776; S66999; S56996; S4
C; Dieckmann, C.L.; Homison, G.; Tzagoloff, A.
A; Dieckmann, C.L.; Homison, G.; Tzagoloff, A.
A; Title: Assembly of the mitochondrial membrane
A; Reference number: S05829; MUID:84185566
A; Accession: S03829
A; Molecular type: DNA
A; Residues: 1-654 </D>
A; Cross-references: EMBL:K02647; NID:9171166; E
R; Vandembroucq, M.; Durand, P.; Boile, P.A.; Dion,
Year 10, 1657-1662, 1994
A; Title: Sequence analysis of a 40.2 kb DNA fra
A; Reference number: S50701; MUID:95242842
A; Accession: S50776
A; Status: nucleic acid sequence not shown; tra
A; Molecular type: DNA
A; Residues: 1-654 </D>

```

A; Cross-references: EMBL:Z34098; PIDN:CAA84002.1; PID:9496953
 A; Note: the nucleotide sequence was submitted to the EMBL Data Library, June 1994
 R; Vanderhol, M.; Durand, P.; Portetelle, D.; Hilger, F.
 submitted to the Protein Sequence Database, September 1995
 A; Reference number: S56835
 A; Accession: S56919
 A; Molecule type: DNA
 A; Residues: 1-654 <VAN>
 A; Cross-references: EMBL:249484; PIDN:CAA89506.1; PID:gi015591; GSPDB:GN00
 R; Purnelle, B.; Coster, F.; Goffeau, A.
 submitted to the Protein Sequence Database, September 1995
 A; Reference number: S56977
 A; Accession: S56936
 A; Molecule type: DNA
 A; Residues: 637-654 <PUR>
 A; Cross-references: EMBL:Z49484; GSPDB:GN00010; MIPS:YJL209w
 C; Genetics:
 A; Gene: SGD:csp1; MIPS:YJL209w
 A; Cross-references: SGD:S0003745; MIPS:YJL209w
 A; Map position: 10L
 A; Genome: nuclear
 C; Function:
 A; Description: pre-mRNA processing
 A; Note: required for correct 5' terminal processing of cytochrome b pre-mRNA
 C; Superfamily: CBF1 protein
 C; Keywords: mitochondrion
 RESULT 3
 Query Match 78.3%; Score 36; DB 1; Length 654;
 Best Local Similarity 75.0%; Pred. No. 44;
 Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 Qy 1 QKLCHOKK 8
 Db 514 KKLCHYKK 521

RESULT 3
 Query Match 78.3%; Score 36; DB 1; Length 654;
 Best Local Similarity 75.0%; Pred. No. 44;
 Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 Qy 1 QKLCHOKK 8
 Db 514 KKLCHYKK 521

RESULT 3
 Query Match 78.3%; Score 36; DB 1; Length 654;
 Best Local Similarity 75.0%; Pred. No. 44;
 Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 Qy 1 QKLCHOKK 8
 Db 514 KKLCHYKK 521

Query Match 73.9%; Score 34; DB 2; Length 1188;
 Best Local Similarity 71.4%; Pred. No. 1.6e+02;
 Matches 2; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 QKLCHOK 7
 Db 1117 BELCHQK 1123

RESULT 5
 T30528
 reverse transcriptase - rice blast fungus
 C; Species: Magnaporthe grisea (rice blast fungus)
 C; Date: 22-Oct-1999 #sequence_revision 22-Oct-1999 #text_change 22-Oct-1999
 C; Accession: T30528
 R; Neyn III, M.A.; Farrall, L.; Valent, B.; Chumley, F.G.; Orbach, M.J.
 submitted to the EMBL Data Library, August 1997
 A; Description: Magnaporthe grisea repeated DNA element MGR583 is a member of the LI
 A; Reference number: Z20845
 A; Accession: T30528
 A; Status: preliminary; translated from GB/EMBL/DDJB
 A; Molecule type: DNA
 A; Residues: 1-1295 <MEY>
 A; Cross-references: EMBL:AF018033; PID:92454622; PIDN:AAB71689.1
 Query Match 73.9%; Score 34; DB 2; Length 1295;
 Best Local Similarity 71.4%; Pred. No. 1.7e+02;
 Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
 Qy 2 KLCQHK 8
 Db 33 ELCHQRK 39

RESULT 6
 T31422
 C-terminal domain-binding protein rAG - rat
 C; Species: Rattus norvegicus (Norway rat)
 C; Date: 29-Oct-1999 #sequence_revision 29-Oct-1999 #text_change 07-Dec-1999
 C; Accession: T31422
 R; Yuryev, A.; Purturajan, M.; Litingtung, Y.; Joshi, R.V.; Gentile, C.; Gebara, M.;
 Proc. Natl. Acad. Sci. U.S.A. 93: 6915-6980, 1996
 A; Title: The C-terminal domain of the largest subunit of RNA polymerase II interact
 A; Reference number: 221024; MUID:96293459
 A; Accession: T31422
 A; Status: preliminary; translated from GB/EMBL/DDJB
 A; Molecule type: mRNA
 A; Residues: 1-1473 <YUR>
 A; Cross-references: EMBL:U9057; PID:gi1438533; PID:91438534; PIDN:AAAC52658.1
 A; Experimental source: hippocampus

Query Match 73.9%; Score 34; DB 2; Length 1473;
 Best Local Similarity 71.4%; Pred. No. 1.9e+02;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 Qy 1 QKLCHOK 7
 Db 427 RKLCHEK 433

RESULT 4
 A71621
 protein with 5'-3' exonuclease domain (Kem-1 family) PFB0205C - malaria parasite (Plasmo
 C; Species: Plasmodium falciparum
 C; Date: 13-Nov-1998 #sequence_revision 13-Nov-1998 #text_change 21-Jul-2000

QY 1 QKICHQK 7
 Db 1425 QKICHSK 1431

RESULT 7
 E72675 hypothetical protein APE0826 - Aeropyrum pernix (strain K1)
 C;Species: Aeropyrum pernix
 C;Date: 20-Aug-1999 #sequence_revision 20-Aug-1999 #text_change 20-Jun-2000
 C;Accession: E72675
 R;Kawarabayasi, Y.; Hino, Y.; Horikawa, H.; Yamazaki, S.; Haikawa, Y.; Jin-no, K.; Takahashi, T.; Tanamiya, M.; Masuda, S.; Furahashi, T.; Tanaka, T.; Kudo, Y.; Yamazaki, J.; Kawamura, H.; Niernberg, W.C.; White, O.; Eisen, J.A.; Salzberg, S.L.; Fraser, C.M.; Venet, A;Title: Complete genome sequence of an aerobic hyper-thermophilic Crenarchaeon, Aeropyrum pernix: Complicated genome sequence of an aerobic hyper-thermophilic Crenarchaeon, Aeropyrum pernix
 A;Reference number: A72450; MUID:99310339
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 1-263 <KAW>
 A;Cross-references: DDBJ:APP00060; NID:g5104188; PIDN:BAA79805.1; PID:g5104490
 A;Experimental source: strain K1
 C;Genetics:
 A;Gene: APE0826
 C;Superfamily: Aeropyrum pernix hypothetical protein APE0826

Query Match 71.7%; Score 33; DB 2; Length 263;
 Best Local Similarity 83.3%; Pred. No. 73; Mismatches 0; Indels 0; Gaps 0;
 Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 3 LCHQKK 8
 Db 258 LCHQKR 263

RESULT 8
 S36336 probable transcription factor PLZF - human
 C;Species: Homo sapiens (man)
 C;Date: 31-Dec-1993 #sequence_revision 31-Dec-1993 #text_change 05-Nov-1999.
 C;Accession: S36336; S31989
 R;Chen, Z.; Brand, N.J.; Chen, A.; Chen, S.J.; Tong, J.-H.; Wang, Z.-Y.; Waxman, S.; Zelenina, N.; White, S.; EMBO J. 12, 1161-1167, 1993
 A;Title: Fusion between a novel Krueppel-like zinc finger gene and the retinoic acid receptor α gene
 A;Reference number: S36336; MUID:9320216
 A;Accession: S36336
 A;Molecule type: mRNA
 A;Residues: 1-73 <CHE>
 A;Cross-references: EMBL:Z19002; NID:g380517; PIDN:CAA79489.1; PID:g38518
 C;Genetics:
 A;Gene: PLZF
 C;Superfamily: POZ domain homology
 F;Keywords: Zinc finger
 F;20-118/Domain: poz domain homology <POZ>

Query Match 71.7%; Score 33; DB 2; Length 673;
 Best Local Similarity 83.3%; Pred. No. 1.5e+02; Mismatches 0; Indels 0; Gaps 0;
 Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 2 KLCHQK 7
 Db 605 KLCHQR 610

RESULT 9
 T00554 hypothetical protein At2g39440 [Imported] - Arabidopsis thaliana
 N;Alternate names: hypothetical protein F12L6.10
 C;Species: Arabidopsis thaliana (mouse-ear cress)
 C;Date: 01-Feb-1999 #sequence_revision 01-Feb-1999 #text_change 16-Feb-2001
 C;Accession: T00554; D84517

Query Match 71.7%; Score 33; DB 2; Length 1247;
 Best Local Similarity 71.4%; Pred. No. 2.5e+02; Mismatches 1; Indels 0; Gaps 0;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 2 KLCHQKK 8
 Db 265 KICHQAK 271

RESULT 11
 W6WIEP E6 Protein - European elk papillomavirus
 C;Species: European elk papillomavirus
 C;Date: 31-Mar-1989 #sequence_revision 31-Mar-1989 #text_change 11-May-2000
 C;Accession: A29499; F94437; F345056

R;Ahola, H.; Bergman, P.; Stroem, A.C.; Moreno-Lopez, J.; Pettersson, U.
 Gene 50, 195-205, 1986
 A;Title: Organization and expression of the transforming region from the European elk p8
 A;Reference number: A91567; MUID:87219878
 A;Accession: A29499
 A;Molecule type: DNA
 A;Residues: 1-135 <AHO>
 A;Cross-references: GB:MI15933; NID:g333025; PIDN:AAA66849.1; PID:9484015
 R;Eriksson, A.
 unpublished results 1987, cited by GenBank
 A;Reference number: A94457
 A;Accession: F94457
 A;Molecule type: DNA
 A;Residues: 1-135 <ERI>
 A;Cross-references: GB:MI15933; NID:g333025; PIDN:AAA66849.1; PID:9484015
 R;Pettersson, U.
 submitted to GenBank, August 1987
 A;Reference number: A94506
 A;Accession: F94506
 A;Molecule type: DNA
 A;Residues: 1-135 <PET>
 A;Cross-references: GB:MI15933; NID:g333025; PIDN:AAA66849.1; PID:9484015
 C;Superfamily: Papillomavirus E6 Protein
 C;Keywords: DNA binding; early protein; zinc finger

Query Match 69.6% Score 32; DB 1; Length 135;
 Best Local Similarity 71.4%; Pred. No. 65;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 2 KLCHQKK 8
 Db 24 KRCHEEKK 30

RESULT 12
 hypothetical protein F21H12.3 - *Caenorhabditis elegans*
 C;Species: *Caenorhabditis elegans*
 C;Date: 20-Sep-1999 #sequence_revision 20-Sep-1999 #text_change 24-Nov-1999
 R;Favelllo, T.
 submitted to the EMBL Data Library, July 1995

A;Description: The sequence of *C. elegans* cosmid F21H12.

A;Reference number: Z1B464
 A;Accession: T16125

A;Status: Preliminary; translated from GB/EMBL/DBJ

A;Residues: 1-210 <FAV>

A;Cross-references: EMBL:U23176; NID:g726404; PID:9726407; PIDN:ACAC46715.1; CEESP:F21H12.3
 A;Experimental source: strain Bristol N2
 C;Genetics:

A;Gene: CEESP:F21H12.3

A;Introns: 37/3; 62/2
 C;Superfamily: *Caenorhabditis elegans* hypothetical protein F21H12.3

Query Match 69.6% Score 32; DB 2; Length 210;
 Best Local Similarity 71.4%; Pred. No. 92;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 QKLCHQK 7
 Db 23 KNLCHQK 29

RESULT 13
 I39522 3-dehydroquinate dehydratase (EC 4.2.1.10), catabolic [validated] - *Acinetobacter calcoaceticus*
 C;Species: *Acinetobacter calcoaceticus*
 C;Accession: I39522
 R;Elsemore, D.A.; Ornston, L.N.

J. Bacteriol. 177, 5971-5978, 1995

A;Title: Unusual ancestry of dehydratases associated with quinate catabolism in *Aci-*

*n*etria

A;Reference number: I39522; MUID:601189

A;Accession: I39522
 A;Status: Preliminary; translated from GB/EMBL/DBJ

A;Molecule type: DNA

A;Residues: 1-240 <RES>

A;Cross-references: EMBL:U20284; PID:9644872; PID:9644873

C;Genetics:

A;Gene: quiB

C;Function:

A;Description: EC 4.4.2.1.10 [validated; MUID:96011389]

C;3-dehydroquinate dehydratase; 3-dehydroquinate dehydratase homology

C;Superfamily: 3-dehydroquinate dehydratase; hydro-lyase

C;Keywords: carbon-oxygen lyase; 3-dehydroquinate dehydratase homology <DQD>

F:38-277/Domain: 3-dehydroquinate dehydratase homology

unpublished results 1987, cited by GenBank

A;Reference number: A94457

A;Accession: F94457

A;Molecule type: DNA

A;Residues: 1-135 <ERI>

A;Cross-references: GB:MI15933; NID:g333025; PIDN:AAA66849.1; PID:9484015

R;Pettersson, U.

submitted to GenBank, August 1987

A;Reference number: A94506

A;Accession: F94506

A;Molecule type: DNA

A;Residues: 1-135 <PET>

A;Cross-references: GB:MI15933; NID:g333025; PIDN:AAA66849.1; PID:9484015

C;Superfamily: Papillomavirus E6 Protein

C;Keywords: DNA binding; early protein; zinc finger

Query Match 69.6% Score 32; DB 1; Length 290;

Best Local Similarity 85.7%; Pred. No. 1.2e+02;

Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 KLCHQKK 8
 Db 167 KLAHQKK 173

RESULT 14
 S26707 transcription initiation factor IIB - yeast (Saccharomyces cerevisiae)

N;Alternate names: protein P9513.4; protein YPRO86W

C;Species: *Saccharomyces cerevisiae*

C;Date: 10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change 21-Jul-2000

C;Accession: S26707; S69072

R;Pinto, I.; Ware, D.E.; Hampsey, M.

Cell 68, 977-988, 1992

A;Title: The yeast SUA7 gene encodes a homolog of human transcription factor TFIIB

A;Reference number: S26707; MUID:92191276

A;Accession: S26707

A;Molecule type: DNA

A;Cross-references: EMBL:M81380; MUID:9172776; PIDN:AAA35126.1; PID:9172777

A;Experimental source: strain T15

R;Couch, J.

submitted to the EMBL Data Library, March 1996

A;Description: The sequence of *S. cerevisiae* cosmid 9513.

A;Reference number: S69072

A;Accession: S69072

A;Molecule type: DNA

A;Residues: 1-345 <COU>

A;Cross-references: EMBL:U51033; NID:91230676; PIDN:AAE68135.1; PID:91230692; GSPDB

C;Genetics:

A;Gene: SGD:SUA7; MIPS:YPRO86W

A;Cross-references: SGD:50006290; MIPS:YPRO86W

A;Map position: 16R

C;Superfamily: transcription initiation factor IIB; transcription initiation factor IIB homology <TFIIB>

F:23-318/Domain: transcription initiation factor IIB homology <TFIIB>

F:24-48/Region: zinc finger CCCC motif

F:133-210,239-313/Region: duplication

Query Match 69.6% Score 32; DB 1; Length 345;

Best Local Similarity 71.4%; Pred. No. 1.4e+02;

Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 2 KLCHQKK 8
 Db 155 KLCHEDEK 161

RESULT 15
 E6233 membrane transport protein (glnQ) homolog MG303 - Mycoplasma genitalium

C;Species: Mycoplasma genitalium
C;Date: 17-Nov-1995 #sequence_revision 17-Nov-1995 #text_change 02-Feb-2001
C;Accession: EG4233
R;Fraser, C.M.; Gocayne, J.D.; White, O.; Adams, M.D.; Clayton, R.A.; Fleischmann, R.D.;
M.; Fuhrtner, J.; Nguyen, D.; Utterback, T.R.; Sudek, D.M.; Phillips, C.A.; Merrick, J.;
C.A.; Venter, J.C.
Science 270, 397-403, 1995
A;Title: The minimal gene complement of Mycoplasma genitalium.
A;Reference number: A64200; MUID: 96026346
A;Accession: EG4233
A;Status: preliminary; nucleic acid sequence not shown; translation not shown
A;Molecule type: DNA
A;Residues: 1-357 <TIGR>
A;Cross-references: GB:U39711; GB:L43967; NID:91045997; PID:g1046002; TIGR:MG303
A;Experimental source: strain G-37
C;Genetics:
A;Generic code: SGC3
C;Superfamily: unassigned ATP-binding cassette proteins; ATP-binding cassette homology
C;Keywords: ATP; nucleotide binding; P-loop
F;90-388/Domain: ATP-binding cassette homology <ABC>
F;107-114/Region: nucleotide-binding motif A (P-loop)

Query	Match	Score	Local	Similarity	Pred.	No.	Length	DB	Gaps	Indels	Mismatches
Qy	2	KLCHQK	7	69.6%	32;	83.3%	357;	2;	0;	0;	0;
Db	183	KLCHHK	188								

Search completed: February 12, 2002, 11:51:40
Job time: 301 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on : February 12, 2002, 11:49:43 ; Search time 98.92 Seconds

Perfect score: 46
Sequence: 1 QKLCHOKK 8

Scoring table: BL2SUM62
Gapop 10.0 , Gapext 0.5

Searched:

522463 seqs, 74073290 residues

Total number of hits satisfying chosen parameters :

522463

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : A_Geneseq_1101;*

1: /SIDSB/gcdata/geneseq/geneseq/AA1980.DAT:*

2: /SIDSB/gcdata/geneseq/geneseq/AA1981.DAT:*

3: /SIDSB/gcdata/geneseq/geneseq/AA1982.DAT:*

4: /SIDSB/gcdata/geneseq/geneseq/AA1984.DAT:*

5: /SIDSB/gcdata/geneseq/geneseq/AA1985.DAT:*

6: /SIDSB/gcdata/geneseq/geneseq/AA1986.DAT:*

7: /SIDSB/gcdata/geneseq/geneseq/AA1987.DAT:*

8: /SIDSB/gcdata/geneseq/geneseq/AA1988.DAT:*

9: /SIDSB/gcdata/geneseq/geneseq/AA1989.DAT:*

10: /SIDSB/gcdata/geneseq/geneseq/AA1990.DAT:*

11: /SIDSB/gcdata/geneseq/geneseq/AA1991.DAT:*

12: /SIDSB/gcdata/geneseq/geneseq/AA1992.DAT:*

13: /SIDSB/gcdata/geneseq/geneseq/AA1993.DAT:*

14: /SIDSB/gcdata/geneseq/geneseq/AA1994.DAT:*

15: /SIDSB/gcdata/geneseq/geneseq/AA1995.DAT:*

16: /SIDSB/gcdata/geneseq/geneseq/AA1996.DAT:*

17: /SIDSB/gcdata/geneseq/geneseq/AA1997.DAT:*

18: /SIDSB/gcdata/geneseq/geneseq/AA1998.DAT:*

19: /SIDSB/gcdata/geneseq/geneseq/AA1999.DAT:*

20: /SIDSB/gcdata/geneseq/geneseq/AA2000.DAT:*

21: /SIDSB/gcdata/geneseq/geneseq/AA2001.DAT:*

22: /SIDSB/gcdata/geneseq/geneseq/AA2002.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query	Match Length	DB ID	Description
1	36	78.3	91	22 AAB75580	Human secreted protein
2	35	76.1	551	20 AA29955	Mouse G1CE protein
3	34	73.9	88	21 AA56915	Human prostate cancer
4	34	73.9	102	22 AAC76544	Human Colon cancer
5	34	73.9	522	22 AA995513	Human protein sequ
6	34	73.9	790	22 AA993045	Human Protein sequ
7	34	73.9	1188	21 AAB18183	Plasmidium falciparum
8	33	71.7	197	21 AAO8675	Arabidopsis thaliana
9	33	71.7	452	21 AA56974	Arabidopsis thaliana
10	33	71.7	678	22 AA25801	Human protein sequ
11	33	71.7	778	21 AA79180	Haemato poetic stellate protein

Wild-type yeast tr					
Loss-of-function m					
Loss-of-function m					
Loss-of-function m					
Loss-of-function m					
Loss-of-function m					
Loss-of-function m					
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Loss-of-function m					
Loss-of-function m					
Peptide #10214 enc					
Peptide #1569 enc					
Peptide #1627 enc					
Peptide #1558 enc					
Amino acid sequenc					
Recombinant human					
Mature rHuFP doma					
Human novel protei					
Recombinant human					
Mature rHuFP doma					
Human secreted pro					
Human novel protei					
Alpha foetoprotein					
Human Alpha-fetop					
Human granulocyte					
Dioxin receptor am					
Human secreted pro					
Human novel protei					
Alpha foetoprotein					
Human colon cancer					
Human Alpha protein comp					
Human granulocyte					
AAU15285					
AAU36177					
AAU99228					
AAU362068					
AAU14513					
AAU99222					
AAU62071					
AAU99227					
AAU62067					
AAU97388					
AAU78169					
AAU14277					
AAU10123					
AAU99222					
AAU71130					
AAU92680					
AAU97388					
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AAU14489					
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AAU14277					
AAU14489					

DR N-PSDB; AAF64203.
 XX Nucleic acid molecules encoding human secreted proteins, used in preventing, treating or ameliorating a disorder, e.g. Alzheimer's and Parkinson's diseases and cancers -
 PT Disclosure; Page 60: 542pp; English.
 XX Human secreted proteins AAB75506 - AAB75554 are encoded by polynucleotide sequences AAF64176 - AAF64224. The specification includes amino acid sequences AAB75555 - AAB75606 which represent fragments of the human secreted proteins, and protein sequences with which they share homology. The proteins and polynucleotides, their agonists and antagonists have activities dependent on the tissues and cells in which they are expressed, examples of these activities include, immunosuppressive; antiarthritic; antiproliferative; cytostatic; cariostatic; vasotropic; cerebroprotective; nortropic; neuroprotective; antibacterial; virucide; fungicide; opthalmological; and vulnerary. The proteins, polynucleotides, agonists and antagonists can be used to treat or detect or diagnose various diseases and disorders including, autoimmune diseases e.g. rheumatoid arthritis, hyperproliferative disorders e.g. neoplasms of the breast or liver, cardiovascular disorders e.g. cardiac arrest, cerebrovascular disorders e.g. cerebral ischaemia, angiogenesis, nervous system disorders e.g. Alzheimer's disease, infections caused by bacteria, viruses and fungi and ocular disorders e.g. corneal infection. The polypeptides can also be used to aid wound healing and epithelial cell proliferation, to prevent skin ageing due to sunburn, to maintain organs before transplantation, for supporting cell culture of primary tissues, to regenerate tissues and in Chemosaxis. The polypeptides can also be used as a food additive or preservative to increase or decrease storage capabilities. Included in the invention are polynucleotide sequences AAF64167 - AAF64175 and peptide AAB75505 which are used in the isolation, identification and characterisation of the proteins of the invention.

XX Sequence 91 AA;
 SQ Query Match 78.3%; Score 36; DB 22; Length 91;
 Best Local Similarity 100.0%; Pred. No. 20;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 QKLCHQ 6
 Db 30 qklichq 35

RESULT 2
 AAY29955 ID AAY29955 standard; Protein: 551 AA.
 XX AC AAY29955;
 XX DT 22-NOV-1999 (first entry)
 DE Mouse CG1CE Protein sequence.
 XX CG1CE: Best's macular dystrophy; mutation; diagnosis; detection; KW CG1CE; Best's macular dystrophy; mutation; diagnosis; detection;
 KW BMD; age-related macular dystrophy.
 XX OS Mus sp.
 PN WO943695-A1.
 XX PD 02-SEP-1999.
 XX PF 22-FEB-1999; 99WO-US03790.
 XX PR 25-FEB-1998; 98US-0075941.
 PR 18-DEC-1998; 98US-012926.
 PA (MERCK & CO INC.
 PA (UYUP-) UNIV UPPSALA.

XX Petrukhin K, Caskey CT, Metzker M, Wadeius C;
 PI XX DR WPI: 1999-540560/45.
 PT XX DR N-PSDB; AAZ2129.
 PS XX PT Human and mouse polynucleotides encoding CG1CE polypeptides -
 XX PS Claim 7; Fig 8; 67pp; English.
 CC The present sequence represents the mouse CG1CE protein. When the CG1CE gene is mutated it is responsible for Best's macular dystrophy (BMD). Polynucleotides encoding CG1CE are useful for diagnosing whether a patient carries a mutation in the CG1CE gene. Normal and mutated CG1CE proteins are useful for identifying activators and/or inhibitors of these proteins, in order to treat BMD. The CG1CE gene offers a simpler and cheaper method of diagnosing BMD without the need for the presence of the patient. The gene may also be useful to discovering the genetic cause of age-related macular dystrophy.
 XX SQ Sequence 551 AA;
 CC Query Match 76.1%; Score 35; DB 20; Length 551;
 Best Local Similarity 62.5%; Pred. No. 1.5e+02;
 Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;
 QY 1 QKLCHQK 8
 Db 490 qeichmkk 497

RESULT 3
 AAB55915 ID AAB55915 standard; Protein: 88 AA.
 XX AC AAB55915;
 XX DT 13-MAR-2001 (first entry)
 DE Human Prostate cancer antigen protein sequence SEQ ID NO:1493.
 XX Human Prostate cancer antigen protein sequence SEQ ID NO:1493.
 XX AC AAB55915;
 XX DT 13-MAR-2001 (first entry)
 DE Human Prostate cancer antigen protein sequence SEQ ID NO:1493.
 XX KW Prostate cancer; prostate cancer antigen; detection; diagnosis;
 KW neuroprotective; cytoactive; immunomodulatory; muscular;
 KW cardioactive; nephrotoxic; antiinfective; gynaecological;
 KW antibacterial; gene therapy; neural; immune; reproductive; renal;
 KW gastrointestinal; pulmonary; cardiovascular; proliferative disorder;
 KW wound; infectious disease.
 XX Homo sapiens.
 OS PN WO200055174-A1.
 XX PD 21-SEP-2000.
 XX PF 08-MAR-2000; 2000WO-US05988.
 XX PR 12-MAR-1999; 99US-0124270.
 PA (HUMA-) HUMAN GENOME SCI INC.
 PA (ROSE/) ROSEN C A.
 PI PN Rosen CA, Ruben SM;
 XX DR 2000-587513/55.
 DR N-PSDB; AAI6118.
 PT Prostate cancer associated gene sequences, referred to as prostate cancer antigens, useful for treatment, prevention, and diagnosis of PT disorders such as prostate cancer -
 XX PT
 PS Claim 11; Page 1930-1931; 2338pp; English.

AAF16566 to AAF16505 encode the human prostate cancer associated proteins, called prostate cancer antigens, given in AAB5663 to AAB57302. The prostate cancer antigens can have neuroprotective, cytostatic, cardioactive, immunomodulatory, muscular, pulmonary, gastrointestinal, nephrotoxic, antiinfective, gynaecological and antibacterial activities, and can be used in gene therapy. The prostate cancer antigen polynucleotides may be used for detection of prostate cancer, chromosome identification, as chromosome markers, and for numerous other diagnostic or research purposes. The prostate cancer antigens may be used to treat disorders such as neural, immune, muscular, reproductive, gastrointestinal, pulmonary, cardiovascular, renal, and proliferative disorders, wounds, and infectious diseases. AAF16506 to AAF16514 to AAB57303 represent sequences used in the exemplification of the present invention.

Sequence 88 AA:
Query Match 73.9%; Score 34; DB 21; Length 88;
Best Local Similarity 62.5%; Pred: No. 44; Indels 0; Gaps 0;
Matches 5; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 QKLCHOKK 8
Db 62 qelchqk 69

RESULT 4
AAG70544
ID AAG70544 standard; Protein; 102 AA.
XX
AC AAG70544;
XX
DT 03-SEP-2001 (first entry)
XX Human colon cancer antigen protein SEQ ID NO:7308.
DE XX
KW Human; colon cancer; colon cancer antigen; diagnosis; detection;
KW colorectal carcinoma.
XX
OS Homo sapiens.
XX
PN WO200122920-A2.
XX
PD 05-APR-2001.
XX
PF 28-SEP-2000; 2000WO-US26524.
XX
PR 29-SEP-1999; 99US-0157137.
PR 03-NOV-1999; 99US-0163380.
PA (HUMA-) HUMAN GENOME SCI INC.
XX
PI Ruben SM, Barash SC, Birse CE, Rosen CA;
XX
DR WPI; 2001-235357/24.
XX
PS DR; AAH35949.

Nucleic acids encoding 4277 human colon cancer-associated polypeptides, useful for preventing, diagnosing and/or treating colorectal cancers -
XX
PT Claim 11; Page 8741-8742; 9803pp; English.
XX
PT CC AH32943 to AAH37195 and AAG73514 to AAG77788 represent human colon cancer-associated nucleic acid molecules (N) and proteins (P), where the proteins are collectively known as colon cancer antigens. The colon cancer antigens have cytostatic activity and can be used in gene therapy and vaccine production. N and P may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate P expression. For example, N and P may be used to treat disorders associated with decreased expression by rectifying mutations or deletions in a patient's genome that affect the activity of P by expressing inactive proteins or to supplement the patients own production of P.

CC Additionally, N may be used to produce the colon cancer-associated PS, by inserting the nucleic acids into a host cell and culturing the cell to express the proteins. N and P can be used in the prevention, diagnosis and treatment of colorectal carcinomas and cancers. AAH37195 to AAH37204 and AAB7789 represent sequences used in the exemplification of the present invention.

CC N.B. Pages 666 to 682 and page 7053 of the sequence listing were missing at time of publication, meaning no sequences are present for SEQ ID NO:1052, 7921 and 7922.

CC SQ Sequence 102 AA;
Query Match 73.9%; Score 34; DB 22; Length 102;
Best Local Similarity 83.3%; Pred. No. 50;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 3 LCHQQKK 8
Db 61 ichqkk 66

RESULT 5
AAB95513
ID AAB95513 standard; Protein; 522 AA.
XX
AC AAB95513;
XX
DT 26-JUN-2001 (first entry)
XX
DE Human protein sequence SEQ ID NO:18081.
XX
KW Human; primer; detection; diagnosis; antisense therapy; gene therapy.
XX
OS Homo sapiens.
XX
PN EPI074617-A2.
XX
PD 07-FEB-2001.
XX
PF 28-JUL-2000; 2000EP-0116126.
XX
PR 29-JUL-1999; 99JP-0248036.
PR 27-AUG-1999; 99JP-0300253.
PR 11-JAN-2000; 2000JP-018776.
PR 02-MAY-2000; 2000JP-0183767.
PR 09-JUN-2000; 2000JP-0241899.
XX
PA (HELI-) HELIX RES INST.
XX
PI Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J,
PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otusuki T;
XX
WPI; 2001-318749/34.

XX
PT Primer sets for synthesizing polynucleotides, particularly the 5602 full-length cDNAs defined in the specification, and for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs -
XX
PT Claim 8; SEQ ID 18081; 2537pp + CD ROM; English.
XX
PT The present invention describes primer sets for synthesising 5602 full-length cDNAs defined in the specification. Where a primer set comprises: (a) an oligo-dT primer and an oligonucleotide complementary to the complementary strand of a polynucleotide which comprises one of the 5602 nucleotide sequences defined in the specification, where the oligonucleotide comprises at least 15 nucleotides; or (b) a combination of an oligonucleotide comprising a sequence complementary to the complementary strand of a polynucleotide which comprises a 5'-end sequence and an oligonucleotide comprising a sequence complementary to a polynucleotide which comprises a 3'-end sequence, where the oligonucleotide comprises at least 15 nucleotides and the combination of

the 5'-end sequence/3'-end sequence is selected from those defined in the specification. The primer sets can be used in antisense therapy and particularly full-length cDNAs. The primers are useful for synthesising polynucleotides, detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs. The primers allow obtaining of the full-length cDNAs easily without any specialised methods. AAB03166 to AAH13628 and AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632 represent oligonucleotides, all of which are used in the exemplification of the present invention.

Sequence 522 AA;

	Query Match	Score	DB	Length
Best Local Similarity	62.5%	22;	22;	522;
Matches	5;	Pred. No.	2.2e+02;	
Conservative	3;	Mismatches	0;	
		Indels	0;	
		Gaps	0;	

Qy 1 QKLHQK 8
|:|||:;
Db 157 qelchqg 164

RESULT 6

ID AAB93045 standard; Protein; 790 AA.
XX

AC AAB93045;

XX DT 26-JUN-2001 (first entry)

XX DE Human protein sequence SEQ ID NO:11834.

XX KW Human; primer; detection; diagnosis; antisense therapy; gene therapy.

XX OS Homo sapiens.

XX PN EP1074617-A2.

XX PD 07-FEB-2001.

XX PF 28-JUL-2000; 2000EP-0116126.

XX PR 29-JUL-1999; 99JP-0248036.

PR 27-AUG-1999; 99JP-030253.

PR 11-JAN-2000; 2000JP-0118776.

PR 02-MAY-2000; 2000JP-0183767.

PR 09-JUN-2000; 2000JP-0241899.

PA (HELI-) HELIX RES INST

XX Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;

PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;

DR 2001-318749/34.

XX PS Claim 8; SEQ ID 11834; 2537PP + CD ROM; English.

XX The present invention describes primer sets for synthesising 5602 full-length cDNAs defined in the specification. Where a primer set comprises: (a) an oligo-dT primer and an oligonucleotide complementary to the complementary strand of a polynucleotide which comprise one of the 5602 nucleotide sequences defined in the specification, where the oligonucleotide comprises at least 15 nucleotides; or (b) a combination of an oligonucleotide comprising a sequence complementary to the complementary strand of a polynucleotide which comprises a 5'-end sequence and an oligonucleotide comprising a sequence complementary to a

polynucleotide which comprises a 3'-end sequence, where the oligonucleotide comprises at least 15 nucleotides and the combination of the 5'-end sequence/3'-end sequence is selected from those defined in the specification. The primer sets can be used in antisense therapy and particularly full-length cDNAs. The primers are useful for synthesising polynucleotides, particularly full-length cDNAs. The primers are also useful for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs. The primers allow obtaining of the full-length cDNAs easily without any specialised methods. AAB03166 to AAH13628 and AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632 represent oligonucleotides, all of which are used in the exemplification of the present invention.

SQ Sequence 790 AA;

	Query Match	Score	DB	Length
Best Local Similarity	62.5%	34;	22;	790;
Matches	5;	Pred. No.	3.2e+02;	
Conservative	3;	Mismatches	0;	
		Indels	0;	
		Gaps	0;	

Qy 1 QKLCHKK 8
|:|||:;
Db 157 qelchqg 164

RESULT 7

ID AAB18183 standard; Protein; 1188 AA.

XX AAB18183;

AC AC

DE 07-NOV-2000 (first entry)

XX Plasmodium falciparum chromosome 2 related protein SEQ ID NO:40.
XX Plasmodium falciparum; chromosome 2; human malaria parasite; vaccine;
KW antimalarial; malaria; protozoacide; infection; insecticide.
XX Plasmodium falciparum.
OS Plasmodium falciparum.
XX WO200025728-A2.
PN XX

PD 11-MAY-2000.

XX PF 05-NOV-1999;

XX PR 05-NOV-1998;

XX PR 99WO-US26796.

XX PR 05-NOV-1998;

XX PR 98US-0107131.

XX PI Hoffman S, Carucci D, Gardner M, Venter JC;

XX DR WPI: 2000-365347/31.

XX PA (HOFF/) HOFFMAN S.
PA (CARU/) CARUCCI D.
PA (GARD/) GARDNER M.
PA (VENT/) VENTER J C.
XX Disclosure: Page 101-104; 577PP; English.

CC The present invention describes proteins and their fragments (I) encoded by chromosome 2 of the human malarial parasite, Plasmodium falciparum. CC Also described are: (1) nucleotide sequences (II) encoding (I); and (2) CC vaccines against P. falciparum infection comprising (I) or (II). CC (1) and (II) are useful for the development of vaccines against CC P. falciparum infection. (I) and polyclonal antisera or a monoclonal CC antibody raised to immunogens comprising the sequences of (II) are CC useful in the detection of infection with P. falciparum. Furthermore, CC (I) (especially when they are rifins or secreted or membrane proteins)

can aid the identification of drugs to treat or prevent *P. falciparum* infection, or they can be used to identify drug resistance and the *P. falciparum*. Sequencing of the Plasmodium chromosome 2 and the subsequent identification of proteins encoded by it will help to expand our understanding of parasite biology, and provide new targets for complexity of the parasitic lifecycle, and provide new targets for vaccine and drug development. Parasite resistance to drugs and mosquito resistance to insecticides have led to resurgence of malaria in many parts of the world, and there is a pressing need for vaccines and new drugs. AAB70078 to AAA70287 and AAB18144 to AAB18352 represent nucleotide and protein sequences given in the present invention, but which are not specifically mentioned within the specification.

Sequence 1188 AA;

Query Match 73.9%; Score 34; DB 21; Length 1188;
 Best Local Similarity 71.4%; Pred. No. 4.6e+02; 2; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QKLCHQK 7
 Db 1117 eelchqk 1123

RESULT 8
 ID AAG08675 standard; Protein: 197 aa.
 XX
 AC AAG08675;
 XX
 DT 17-OCT-2000 (first entry)

DE Arabidopsis thaliana protein fragment SEQ ID NO: 6307.
 XX Protein identification; signal transduction pathway; metabolic pathway;
 KW hybridisation assay; genetic mapping; gene expression control; promoter;
 KW termination sequence.
 XX OS Arabidopsis thaliana.
 XX PN EP1033405-A2.
 XX PD 06-SEP-2000.
 XX PF 25-FEB-2000; 2000EP-0301439.

XX 25-FEB-1999; 99US-0121825.
 PR 05-MAR-1999; 99US-0123110.
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 PR 23-MAR-1999; 99US-0125788.
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 PR 18-AUG-1999; 99US-0149426.
 PR 20-AUG-1999; 99US-0149722.
 PR 20-AUG-1999; 99US-0149723.
 PR 23-AUG-1999; 99US-0149929.
 PR 23-AUG-1999; 99US-0149902.
 PR 25-AUG-1999; 99US-0150566.
 PR 26-AUG-1999; 99US-0150884.
 PR 27-AUG-1999; 99US-0151065.
 PR 27-AUG-1999; 99US-0151066.
 PR 27-AUG-1999; 99US-0151080.
 PR 30-AUG-1999; 99US-0151303.
 PR 31-AUG-1999; 99US-0151438.
 PR 01-SEP-1999; 99US-0151930.
 PR 07-SEP-1999; 99US-0152363.
 PR 10-SEP-1999; 99US-0153070.
 PR 13-SEP-1999; 99US-0153758.
 PR 15-SEP-1999; 99US-0154018.
 PR 16-SEP-1999; 99US-0154039.
 PR 20-SEP-1999; 99US-0154779.
 PR 22-SEP-1999; 99US-0155139.
 PR 23-SEP-1999; 99US-0155486.
 PR 24-SEP-1999; 99US-0155659.
 PR 28-SEP-1999; 99US-0156458.
 PR 29-SEP-1999; 99US-0156596.
 PR 04-OCT-1999; 99US-0157117.
 PR 05-OCT-1999; 99US-0157153.
 PR 06-OCT-1999; 99US-0157865.
 PR 07-OCT-1999; 99US-0158029.
 PR 08-OCT-1999; 99US-0158232.
 PR 12-OCT-1999; 99US-0158369.
 PR 13-OCT-1999; 99US-0159293.
 PR 13-OCT-1999; 99US-0159294.
 PR 13-OCT-1999; 99US-0159295.
 PR 14-OCT-1999; 99US-0159329.
 PR 14-OCT-1999; 99US-0159330.
 PR 14-OCT-1999; 99US-0159331.
 PR 14-OCT-1999; 99US-0159637.
 PR 14-OCT-1999; 99US-0159638.
 PR 18-OCT-1999; 99US-0159584.
 PR 21-OCT-1999; 99US-0160741.
 PR 21-OCT-1999; 99US-0160767.
 PR 21-OCT-1999; 99US-0160768.
 PR 21-OCT-1999; 99US-0160770.
 PR 21-OCT-1999; 99US-0160814.
 PR 21-OCT-1999; 99US-0160815.
 PR 22-OCT-1999; 99US-0160980.
 PR 22-OCT-1999; 99US-0160981.
 PR 22-OCT-1999; 99US-0160989.
 PR 25-OCT-1999; 99US-0161404.
 PR 25-OCT-1999; 99US-0161405.

PR 25-OCT-1999; 99US-0161406.
 PR 26-OCT-1999; 99US-0161359.
 PR 26-OCT-1999; 99US-0161360.
 PR 28-OCT-1999; 99US-0161361.
 PR 28-OCT-1999; 99US-0161920.
 PR 28-OCT-1999; 99US-0161992.
 PR 29-OCT-1999; 99US-0161993.
 PR 29-OCT-1999; 99US-0162142.

Query Match Score 33; DB 21; Length 197;
 Best Local Similarity 71.7%; Pred: No. 1.4e+02;
 Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 2 KLCHOKK 8
 Db 185 Klichkr 191

RESUL 9
 AAG05674
 ID AAG05674 standard; protein; 242 AA.
 AC AAG05674;
 DT 17-OCT-2000 (first entry)
 XX Arabidopsis thaliana protein fragment SEQ ID NO: 6306.
 KW Protein identification; signal transduction pathway; metabolic pathway;
 KW hybridisation assay; genetic mapping; gene expression control; promoter;
 KW termination sequence.
 XX Arabidopsis thaliana.
 OS Arabidopsis thaliana.
 PN EP1033405 A2.
 XX
 PD 06-SEP-2000.
 XX
 PF 25-FEB-2000; 2000EP-0301439.
 XX
 PR 25-FEB-1999; 99US-0121825.
 PR 05-MAR-1999; 99US-0122180.
 PR 09-MAR-1999; 99US-0123548.
 PR 23-MAR-1999; 99US-0125788.
 PR 25-MAR-1999; 99US-0126264.
 PR 29-MAR-1999; 99US-0126785.
 PR 01-APR-1999; 99US-0127462.
 PR 06-APR-1999; 99US-0128734.
 PR 08-APR-1999; 99US-0128714.
 PR 16-APR-1999; 99US-0128845.
 PR 19-APR-1999; 99US-0130077.
 PR 21-APR-1999; 99US-0130449.
 PR 23-APR-1999; 99US-0130891.
 PR 28-APR-1999; 99US-0131449.
 PR 30-APR-1999; 99US-0132408.
 PR 04-MAY-1999; 99US-0132407.
 PR 05-MAY-1999; 99US-0130510.
 PR 06-MAY-1999; 99US-0132486.
 PR 07-MAY-1999; 99US-0132487.
 PR 11-MAY-1999; 99US-0134256.
 PR 14-MAY-1999; 99US-0134218.
 PR 14-MAY-1999; 99US-0134219.
 PR 14-MAY-1999; 99US-0134221.
 PR 14-MAY-1999; 99US-0134221.
 PR 18-MAY-1999; 99US-013470.
 PR 18-MAY-1999; 99US-0134768.
 PR 19-MAY-1999; 99US-0134941.
 PR 20-MAY-1999; 99US-0135124.
 PR 21-MAY-1999; 99US-0133353.
 PR 24-MAY-1999; 99US-0133629.
 PR 25-MAY-1999; 99US-0136021.

PR	27-MAY-1999;	99US-0136392;	04-AUG-1999;	99US-0147204.
PR	28-MAY-1999;	99US-0136782;	04-AUG-1999;	99US-0147302.
PR	01-JUN-1999;	99US-0137222;	05-AUG-1999;	99US-0147192.
PR	03-JUN-1999;	99US-0137528;	05-AUG-1999;	99US-0147260.
PR	04-JUN-1999;	99US-0137502;	06-AUG-1999;	99US-0147303.
PR	07-JUN-1999;	99US-0137724;	06-AUG-1999;	99US-0147416.
PR	08-JUN-1999;	99US-0138094;	09-AUG-1999;	99US-0147193.
PR	10-JUN-1999;	99US-0138540;	09-AUG-1999;	99US-0147935.
PR	10-JUN-1999;	99US-0138847;	10-AUG-1999;	99US-0148171.
PR	14-JUN-1999;	99US-0139119;	11-AUG-1999;	99US-0148319.
PR	16-JUN-1999;	99US-0139452;	12-AUG-1999;	99US-0148441.
PR	16-JUN-1999;	99US-0139453;	13-AUG-1999;	99US-0148565.
PR	17-JUN-1999;	99US-0139459;	13-AUG-1999;	99US-0148684.
PR	18-JUN-1999;	99US-0139492;	16-AUG-1999;	99US-0149468.
PR	18-JUN-1999;	99US-0139454;	17-AUG-1999;	99US-0149175.
PR	18-JUN-1999;	99US-0139455;	18-AUG-1999;	99US-0149426.
PR	18-JUN-1999;	99US-0139456;	20-AUG-1999;	99US-0149722.
PR	18-JUN-1999;	99US-0139457;	20-AUG-1999;	99US-0149723.
PR	18-JUN-1999;	99US-0139458;	20-AUG-1999;	99US-0149329.
PR	18-JUN-1999;	99US-0139459;	23-AUG-1999;	99US-0149902.
PR	18-JUN-1999;	99US-0139460;	23-AUG-1999;	99US-0149320.
PR	18-JUN-1999;	99US-0139461;	25-AUG-1999;	99US-0150566.
PR	18-JUN-1999;	99US-0139462;	25-AUG-1999;	99US-0150884.
PR	18-JUN-1999;	99US-0139463;	26-AUG-1999;	99US-0150884.
PR	18-JUN-1999;	99US-0139750;	27-AUG-1999;	99US-0151165.
PR	18-JUN-1999;	99US-0139763;	27-AUG-1999;	99US-0151066.
PR	21-JUN-1999;	99US-0139817;	27-AUG-1999;	99US-0151080.
PR	22-JUN-1999;	99US-0139899;	30-AUG-1999;	99US-0151303.
PR	23-JUN-1999;	99US-0140353;	31-AUG-1999;	99US-0151438.
PR	23-JUN-1999;	99US-0140354;	01-SEP-1999;	99US-0151930.
PR	24-JUN-1999;	99US-0142055;	07-SEP-1999;	99US-0153363.
PR	28-JUN-1999;	99US-0142095;	10-SEP-1999;	99US-0153407.
PR	29-JUN-1999;	99US-0142390;	13-SEP-1999;	99US-0153758.
PR	08-JUL-1999;	99US-0140991;	15-SEP-1999;	99US-0154018.
PR	09-JUL-1999;	99US-0141287;	16-SEP-1999;	99US-0154039.
PR	01-JUL-1999;	99US-0141842;	20-SEP-1999;	99US-0154779.
PR	01-JUL-1999;	99US-0142154;	22-SEP-1999;	99US-0155139.
PR	06-JUL-1999;	99US-0142055;	23-SEP-1999;	99US-015546.
PR	08-JUL-1999;	99US-0142390;	24-SEP-1999;	99US-0155659.
PR	12-JUL-1999;	99US-0142920;	28-SEP-1999;	99US-0156458.
PR	13-JUL-1999;	99US-0142977;	29-SEP-1999;	99US-0155956.
PR	15-JUL-1999;	99US-0143542;	04-OCT-1999;	99US-0157117.
PR	15-JUL-1999;	99US-0143624;	05-OCT-1999;	99US-0157753.
PR	16-JUL-1999;	99US-0144005;	06-OCT-1999;	99US-0157865.
PR	16-JUL-1999;	99US-0144085;	07-OCT-1999;	99US-0158029.
PR	19-JUL-1999;	99US-0144086;	08-OCT-1999;	99US-0158232.
PR	19-JUL-1999;	99US-0144325;	12-OCT-1999;	99US-0158269.
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PR	19-JUL-1999;	99US-0144332;	13-OCT-1999;	99US-0159584.
PR	21-JUL-1999;	99US-0144333;	21-OCT-1999;	99US-0160741.
PR	21-JUL-1999;	99US-0144334;	22-OCT-1999;	99US-0160980.
PR	22-JUL-1999;	99US-0144335;	22-OCT-1999;	99US-0160768.
PR	20-JUL-1999;	99US-0144352;	21-OCT-1999;	99US-0160776.
PR	22-JUL-1999;	99US-0144352;	25-OCT-1999;	99US-0161404.
PR	21-JUL-1999;	99US-0144884;	21-OCT-1999;	99US-0161405.
PR	22-JUL-1999;	99US-0145089;	25-OCT-1999;	99US-0161406.
PR	21-JUL-1999;	99US-0145192;	26-OCT-1999;	99US-0161359.
PR	23-JUL-1999;	99US-0145086;	26-OCT-1999;	99US-0161360.
PR	23-JUL-1999;	99US-0145145;	26-OCT-1999;	99US-0161361.
PR	23-JUL-1999;	99US-0145088;	28-OCT-1999;	99US-0161920.
PR	02-AUG-1999;	99US-0145218;	28-OCT-1999;	99US-0161992.
PR	02-AUG-1999;	99US-0145224;	28-OCT-1999;	99US-0161993.

PR 29-OCT-1999; 99US-0162142.

Query Match Score 33; DB 21; Length 24;
Best Local Similarity 71.7%; Pred. No. 1.6e+02;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 2 KICHQKK 8
| | | :
Db 230 kichhkr 236

RESULT 10
ARM25801 ID AAM25801 standard; Protein; 678 AA.
XX AC AAM25801;
XX DT 16-OCT-2001 (first entry)
XX DE Human protein sequence SEQ ID NO:1316.
XX DEI
XX Human; cancer; ulcer; HIV infection; human immunodeficiency virus;
XX antiinflammatory; antirheumatic; antiarthritic; immunosuppressive;
XX antibiotic; endocrine; cardiot; central nervous system; virucide;
XX anti-HIV; fungicide; antimutagen; cardiovacular; antianæmic; anaemia;
XX antiaggregant; haemostatic; vuninary; antiulcer; osteopathic; eczema;
XX dermatological; antiallergic; antiasthmatic; antidiabetic; cytotoxic;
XX neuroprotective; antidepressant; nootropic; antiparkinsonian; infection;
XX immunomodulant; gene therapy; antisense therapy; vaccine; inflammation;
XX antiangiolytic; rheumatoid; arthritis; septic shock; pancreatitis;
XX cardiac dysfunction; neuropathology; cariac anaplyaxis; autoimmunity;
XX genetic disease; haematopoietic disorder; platelet disorder; asthma;
XX thrombocytopenia; osteoporosis; severe combined immunodeficiency;
XX allergic rhinitis; diabetes; multiple sclerosis; depression;
XX Alzheimer's disease; Parkinson's disease; neurodegenerative disorder;
XX neurological disorder.

OS Homo sapiens.
XX WO200153455-A2.
XX 26-JUL-2001.
XX PPF 22-DEC-2000; 2000WO-US35017.
XX PR 23-DEC-1999; 99US-0471275.
XX 21-JAN-2000; 2000US-048875.
XX 25-APR-2000; 2000US-0552317.
XX PA (HYSEQ) HYSEQ INC.
XX PI Tang YT, Liu C, Drmanac RT;
XX DR N-PSDB, AAH99742.
XX PT Isolated human polynucleotides encoding polypeptides, useful for the
XX treatment and diagnosis of e.g. cancer, ulcers and HIV infection -
XX Claim 20; Page 274; 1217pp; English.
XX AAC99166 to AAH99904 encode the human proteins given in AAM25225 to
XX AAM25933. The proteins can have activities based on the tissues and
XX cells they are expressed in, such as: antiinflammatory; antirheumatic;
XX antiarthritic; immunosuppressive; antibacterial; endocrine; cardiot;
XX central nervous system; virucide; anti-HIV; cytotoxic; antimutagen;
XX cardiovascular; antianæmic; antiangiolytic; haemostatic; vulnary;
XX antiulcer; osteopathic; dermatological; antiasthmatic; antidiabetic;
XX anticarckinsonian; cytosatic; neuroprotective; antidepressant; nootropic;
XX CC encoding them can be used in gene therapy, antisense therapy and vaccine
CC production. The proteins and polynucleotides are useful for screening for
CC agonists or antagonists of a protein or for the treatment and diagnosis
CC of disorders associated with the activity of a protein e.g. inflammation,
CC rheumatoid arthritis, septic shock, panoreatitis, cardiac dysfunction,
CC neuropathology, cardiac anaphylaxis, viral, bacterial, HIV and fungal
CC infections, autoimmunity, genetic diseases, haematopoietic disorders,
CC anaemia, platelet disorders, thrombocytopaenia, wounds, burns, ulcers,
CC osteoporosis, severe combined immunodeficiency, eczema, allergic
CC rhinitis, asthma, diabetes, cancer, multiple sclerosis, depression,
CC Alzheimer's disease, Parkinson's disease, neurodegenerative and
CC neurological disorders.

SQ Sequence 678 AA;

Query Match Score 33; DB 22; Length 678;
Best Local Similarity 83.3%; Pred. No. 4.1e+02;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 2 KLCHOK 7
Db 610 Klichqr 615

RESULT 11
AAY79180 ID AAY79180 standard; Protein; 778 AA.
XX AC AAY79180;
XX DT 19-JUN-2000 (first entry)
XX DE Haematopoietic stem cell specific protein.
XX XX Haematopoietic stem cell; immune system disorder;
XX KW leukaemia; antileukemic; immunomodulator; therapy; mouse.
XX OS Mus musculus.
XX FH Key Location/Qualifiers
XX FT Misc-difference 9
XX FT /note= "encoded by CAS"
XX FT Misc-difference 21
XX FT /note= "encoded by ASA"
XX FT Misc-difference 32
XX FT /note= "encoded by SAA"
XX FT Misc-difference 738
XX /note= "encoded by TTN"
XX PN WO200011168-A2.
XX PD 02-MAR-2000.
XX PF 20-AUG-1999;
XX PR 21-AUG-1998;
XX PA (UYPFR) UNIV PRINCETON.
XX PI Lemischka I, Moore K;
XX DR WPI: 2000-237650/20.
XX N-PSDB; AAZ94121.

XX The present sequence is that of a mouse haematopoietic stem cell.
XX Hematopoietic stem cell signaling proteins modulating replication and
XX differentiation for treating immune system disorders and leukaemia -
XX Claim 21; Page 221-223; 256pp; English.

XX The present sequence is that of a mouse haematopoietic stem cell.
XX (HSC) specific proteins can have activities based on the tissues and
XX cells they are expressed in, such as: antiinflammatory; antirheumatic;
XX antiarthritic; immunosuppressive; antibacterial; endocrine; cardiot;
XX central nervous system; virucide; anti-HIV; cytotoxic; antimutagen;
XX cardiovascular; antianæmic; antiangiolytic; haemostatic; vulnary;
XX antiulcer; osteopathic; dermatological; antiasthmatic; antidiabetic;
XX anticarckinsonian; cytosatic; neuroprotective; antidepressant; nootropic;
XX CC encoding them can be used in gene therapy, antisense therapy and vaccine
CC production. The proteins and polynucleotides are useful for screening for
CC agonists or antagonists of a protein or for the treatment and diagnosis
CC of disorders associated with the activity of a protein e.g. inflammation,
CC rheumatoid arthritis, septic shock, panoreatitis, cardiac dysfunction,
CC neuropathology, cardiac anaphylaxis, viral, bacterial, HIV and fungal
CC infections, autoimmunity, genetic diseases, haematopoietic disorders,
CC anaemia, platelet disorders, thrombocytopaenia, wounds, burns, ulcers,
CC osteoporosis, severe combined immunodeficiency, eczema, allergic
CC rhinitis, asthma, diabetes, cancer, multiple sclerosis, depression,
CC Alzheimer's disease, Parkinson's disease, neurodegenerative and
CC neurological disorders.

CC transcription factors, splicing factors, capping factors, transport proteins, translation factors or replication factors that modulate HSC activity, especially differentiation or replication. The invention provides claimed methods: for identifying PHS-C-specific nucleic acids; for generating a stem cell/progenitor cell from PHS-C; for identifying the presence of a PHS-C in a sample; for identifying the presence in a sample of a compound that modulates HSC activity, for using such a compound to treat an immune system condition, especially leukemia; for introducing exogenous nucleic acid into a HSC; and for ex vivo expansion of HSCs. Also claimed are vectors, host cells, and an antibody that specifically binds a HSC-specific protein.

XX Sequence 778 AA;

Query Match 71.7%; Score 33; DB 21; Length 778;
Best Local Similarity 62.5%; Pred. No. 4.7e+02;
Matches 5; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

Qy 1 QKLCHQQK 8
Db 736 qsxchekk 743

RESULT 12
AAW79501

ID AAW79501 standard; Peptide: 23 AA.

XX
AC AAW79501;

XX DT 17-DEC-1998 (first entry)

XX Wild-type yeast transcription factor IIB (region 144-160).

XX Yeast transcription factor IIB; YTFIIB; cell growth; mutation; hybrid;

XX human; antifungal drug

XX Saccharomyces cerevisiae.

XX WO9839355-A1.

XX PR 11-SEP-1998.

XX PT 13-JUN-1997; 97WO-US10404.

XX PR 06-MAR-1997; 97US-0812175.

XX PA (CHIL-) CHILDREN'S HOSPITAL MEDICAL CENT.

XX PI Carson DJ, Dorsey MJ, Ma J, Shaw S, Wingfield J;

XX DR 1998-495788/42.

XX Use of peptide fragments of yeast Transcription Factor IIB - to

XX identify compounds that inhibit fungal cell growth, especially to

XX screen for antifungal drugs active against *Candida albicans*

XX Disclosure: Figure 5; 32pp; English.

XX DR 1998-495788/42.

XX Use of Peptide fragments of Yeast Transcription Factor IIB - to

XX identify Compounds that inhibit fungal cell growth, especially to

XX screen for antifungal drugs active against *Candida albicans*

PS Disclosure: Figure 5; 32pp; English.

XX The present sequence represents the wild-type species-specific region of

CC the yeast transcription factor IIB (YTFIIB), amino acid residues 144-166.

CC This sequence has been found to contain 4 amino acids that are vital for

CC cell growth and in vivo activity of YTFIIB, mutations of these residues

CC results in cells having severe growth defects. By the analysis of hybrid

CC molecules in yeast cells, the 4 amino acids that confer yeast specificity

CC were identified as lysine 147, cysteine 149, lysine 151, and glutamic

CC acid 152. This wild-type sequence was used to generate loss-of-function

CC mutants by changing a number of these critical amino acids to the

CC corresponding human residues. The YTFIIB fragment containing the vital 4

CC amino acid residues provides a yeast-specific target for screening

CC libraries to identify new antifungal drugs.

XX

SQ Sequence 23 AA;

CC proteins, splicing factors, capping factors, transport

CC factors or replication factors that modulate

CC HSC activity, especially differentiation or replication. The

CC invention provides claimed methods: for identifying PHS-C-specific

CC nucleic acids; for generating a stem cell/progenitor cell from

CC PHS-C; for identifying the presence of a PHS-C in a sample; for

CC identifying the presence in a sample of a compound that modulates

CC HSC activity, for using such a compound to treat an immune system

CC condition, especially leukemia; for introducing exogenous nucleic

CC acid into a HSC; and for ex vivo expansion of HSCs. Also claimed

CC are vectors, host cells, and an antibody that specifically binds a

CC HSC-specific protein.

XX

SQ Sequence 23 AA;

CC transcription factors, splicing factors, capping factors, transport

CC factors or replication factors that modulate

CC HSC activity, especially differentiation or replication. The

CC invention provides claimed methods: for identifying PHS-C-specific

CC nucleic acids; for generating a stem cell/progenitor cell from

CC PHS-C; for identifying the presence of a PHS-C in a sample; for

CC identifying the presence in a sample of a compound that modulates

CC HSC activity, for using such a compound to treat an immune system

CC condition, especially leukemia; for introducing exogenous nucleic

CC acid into a HSC; and for ex vivo expansion of HSCs. Also claimed

CC are vectors, host cells, and an antibody that specifically binds a

CC HSC-specific protein.

XX

SQ Sequence 23 AA;

CC transcription factors, splicing factors, capping factors, transport

CC factors or replication factors that modulate

CC HSC activity, especially differentiation or replication. The

CC invention provides claimed methods: for identifying PHS-C-specific

CC nucleic acids; for generating a stem cell/progenitor cell from

CC PHS-C; for identifying the presence of a PHS-C in a sample; for

CC identifying the presence in a sample of a compound that modulates

CC HSC activity, for using such a compound to treat an immune system

CC condition, especially leukemia; for introducing exogenous nucleic

CC acid into a HSC; and for ex vivo expansion of HSCs. Also claimed

CC are vectors, host cells, and an antibody that specifically binds a

CC HSC-specific protein.

XX

SQ Sequence 23 AA;

CC transcription factors, splicing factors, capping factors, transport

CC factors or replication factors that modulate

CC HSC activity, especially differentiation or replication. The

CC invention provides claimed methods: for identifying PHS-C-specific

CC nucleic acids; for generating a stem cell/progenitor cell from

CC PHS-C; for identifying the presence of a PHS-C in a sample; for

CC identifying the presence in a sample of a compound that modulates

CC HSC activity, for using such a compound to treat an immune system

CC condition, especially leukemia; for introducing exogenous nucleic

CC acid into a HSC; and for ex vivo expansion of HSCs. Also claimed

CC are vectors, host cells, and an antibody that specifically binds a

CC HSC-specific protein.

XX

SQ Sequence 23 AA;

CC transcription factors, splicing factors, capping factors, transport

CC factors or replication factors that modulate

CC HSC activity, especially differentiation or replication. The

CC invention provides claimed methods: for identifying PHS-C-specific

CC nucleic acids; for generating a stem cell/progenitor cell from

CC PHS-C; for identifying the presence of a PHS-C in a sample; for

CC identifying the presence in a sample of a compound that modulates

CC HSC activity, for using such a compound to treat an immune system

CC condition, especially leukemia; for introducing exogenous nucleic

CC acid into a HSC; and for ex vivo expansion of HSCs. Also claimed

CC are vectors, host cells, and an antibody that specifically binds a

CC HSC-specific protein.

XX

SQ Sequence 23 AA;

CC transcription factors, splicing factors, capping factors, transport

CC factors or replication factors that modulate

CC HSC activity, especially differentiation or replication. The

CC invention provides claimed methods: for identifying PHS-C-specific

CC nucleic acids; for generating a stem cell/progenitor cell from

CC PHS-C; for identifying the presence of a PHS-C in a sample; for

CC identifying the presence in a sample of a compound that modulates

CC HSC activity, for using such a compound to treat an immune system

CC condition, especially leukemia; for introducing exogenous nucleic

CC acid into a HSC; and for ex vivo expansion of HSCs. Also claimed

CC are vectors, host cells, and an antibody that specifically binds a

CC HSC-specific protein.

XX

SQ Sequence 23 AA;

CC transcription factors, splicing factors, capping factors, transport

CC factors or replication factors that modulate

CC HSC activity, especially differentiation or replication. The

CC invention provides claimed methods: for identifying PHS-C-specific

CC nucleic acids; for generating a stem cell/progenitor cell from

CC PHS-C; for identifying the presence of a PHS-C in a sample; for

CC identifying the presence in a sample of a compound that modulates

CC HSC activity, for using such a compound to treat an immune system

CC condition, especially leukemia; for introducing exogenous nucleic

CC acid into a HSC; and for ex vivo expansion of HSCs. Also claimed

CC are vectors, host cells, and an antibody that specifically binds a

CC HSC-specific protein.

XX

SQ Sequence 23 AA;

CC transcription factors, splicing factors, capping factors, transport

CC factors or replication factors that modulate

CC HSC activity, especially differentiation or replication. The

CC invention provides claimed methods: for identifying PHS-C-specific

CC nucleic acids; for generating a stem cell/progenitor cell from

CC PHS-C; for identifying the presence of a PHS-C in a sample; for

CC identifying the presence in a sample of a compound that modulates

CC HSC activity, for using such a compound to treat an immune system

CC condition, especially leukemia; for introducing exogenous nucleic

CC acid into a HSC; and for ex vivo expansion of HSCs. Also claimed

CC are vectors, host cells, and an antibody that specifically binds a

CC HSC-specific protein.

XX

SQ Sequence 23 AA;

CC transcription factors, splicing factors, capping factors, transport

CC factors or replication factors that modulate

CC HSC activity, especially differentiation or replication. The

CC invention provides claimed methods: for identifying PHS-C-specific

CC nucleic acids; for generating a stem cell/progenitor cell from

CC PHS-C; for identifying the presence of a PHS-C in a sample; for

CC identifying the presence in a sample of a compound that modulates

CC HSC activity, for using such a compound to treat an immune system

CC condition, especially leukemia; for introducing exogenous nucleic

CC acid into a HSC; and for ex vivo expansion of HSCs. Also claimed

CC are vectors, host cells, and an antibody that specifically binds a

CC HSC-specific protein.

XX

SQ Sequence 23 AA;

CC transcription factors, splicing factors, capping factors, transport

CC factors or replication factors that modulate

CC HSC activity, especially differentiation or replication. The

CC invention provides claimed methods: for identifying PHS-C-specific

CC nucleic acids; for generating a stem cell/progenitor cell from

CC PHS-C; for identifying the presence of a PHS-C in a sample; for

CC identifying the presence in a sample of a compound that modulates

CC HSC activity, for using such a compound to treat an immune system

CC condition, especially leukemia; for introducing exogenous nucleic

CC acid into a HSC; and for ex vivo expansion of HSCs. Also claimed

CC are vectors, host cells, and an antibody that specifically binds a

CC HSC-specific protein.

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SQ Sequence 23 AA;

CC transcription factors, splicing factors, capping factors, transport

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